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(54) Title: HUMAN NK-3 RELATED PROSTATE SPECIFIC GENE-1		
(57) Abstract The present invention relates to a novel member of the NK family of homeobox genes. In particular, isolated nucleic acid molecules are provided encoding the human NK-3 prostate specific gene 1 (NKX3.1) protein. NKX3.1 polypeptides are also provided as are vectors, host cells and recombinant methods for producing the same. The invention further relates to screening methods for identifying agonists and antagonists of NKX3.1 activity. Also provided are diagnostic methods for detecting prostate cancer and other cancers and therapeutic methods for prostate cancer and other cancers.		

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Human NK-3 Related Prostate Specific Gene-1

Background of the Invention

Field of the Invention

5 The present invention relates to a novel member of the NK family of homeobox genes. More specifically, isolated nucleic acid molecules are provided encoding a human NK-3 related prostate specific gene (NKX3.1). NKX3.1 polypeptides are also provided, as are vectors, host cells and recombinant methods for producing the same. The invention further relates to screening methods for
10 identifying agonists and antagonists of NKX3.1 activity. Also provided are diagnostic methods for detecting prostate cancer and other cancers and therapeutic methods for prostate cancer and other cancers.

Related Art

15 The discovery of the homeobox as a conserved DNA sequence element in several *Drosophila* genes responsible for controlling the identity of body segments prompted searches for related genes in other organisms. Homeoboxes have since been discovered in the genome of all metazoan organisms, and several hundred unique homeobox genes have been defined in mice and humans (Gehring, W.J. *et al.*, *Annu. Rev. Biochem.* 63:487-526 (1994); Stein, S. *et al.*, *Mech. Develop.* 55:91-108 (1996)). The homeobox encodes a 60-amino acid domain, termed the
20 homeodomain, that includes a helix-turn-helix motif recognized to be structurally-related to the DNA binding domain of several procaryotic proteins and to the products of the yeast mating type focus (Laughon, A. and Scott, M.P., *Nature* 310:25-31 (1984); Shepherd, J.C.W. *et al.*, *Nature* 310:70-71 (1984)). NMR and
25 crystallographic analyses have confirmed that the homeodomain binds DNA (Kissinger, C.R. *et al.*, *Cell* 63:579-590 (1990); Otting, G. *et al.*, *EMBO J.*

9:3085-3092 (1990)). As predicted by the nature of the phenotypes produced when these genes are mutated, both biochemical and genetic analyses have established that the products of homeobox genes are transcriptional regulatory molecules (McGinnis, W. and Krumlauf, R., *Cell* 68:283-302 (1992)).

5 The predicted amino acid sequence of the known homeodomains serves as the principal identifier that allows them to be classified into a minimum of 20 distinct groups (Gehring, W.J. *et al.*, *Annu. Rev. Biochem.* 63:487-526 (1994); Stein, S. *et al.*, *Mech. Develop.* 55:91-108 (1996)). The NK family of homeobox genes, first defined by four related *Drosophila* genes, NK-1 through NK-4, can be
10 separated into two distinct classes. NK-2, -3 and -4 are more related to each other than to other homeobox genes, whereas NK-1 is a more distant relative (Kim, Y. and Nirenberg, M., *Proc. Natl. Acad. Sci. USA* 86:7716-7720 (1989)). In mouse, six NK-2-like genes have been identified (Price, M. *et al.*, *Neuron* 8:241-255 (1992); Lints, T.J. *et al.*, *Development* 119:419-431 (1993)). Three
15 of these are more related to NK-2 than the others, which may themselves form a distinct subclass (Lints, T.J. *et al.*, *Development* 119:419-431 (1993)).

 The majority of studies aimed at characterizing the functions of homeobox genes have focused principally on their developmental roles (McGinnis, W. and Krumlauf, R., *Cell* 68:283-302 91992); Krumlauf, R., *Cell* 78:191-201 (1994)).
20 A prominent example is the Hox family of genes, whose members have been demonstrated to play critical roles in pattern formation during embryogenesis along the anteroposterior body axis of divergent species (Krumlauf, R., *Cell* 78:191-201 (1994)). Some of the Hox genes, as well as members of other classes of homeobox genes, are also expressed during organogenesis, and a few of these
25 have been reported to be expressed in adult tissues. Surprisingly, the potential roles of homeobox genes in fully differentiated tissues and organs have received comparatively little attention. However, the need for patterning functions to maintain the differentiated states of cell populations and to direct the renewal of specific cell types in adults is axiomatic.

The mechanisms involved in the development and maintenance of prostatic tissue are poorly understood. Although it has been recognized for years that normal development and continued expression in adults of the male secondary sexual phenotype is androgen-dependent, there is relatively little known about the genes on which androgens act or the downstream pathways that lead to development of differentiated tissue. As with prostate development, the fundamental mechanisms underlying prostate cancer also remain obscure, however, androgen regulation and the loss thereof plays a critical role. In both developing and mature prostate, the maintenance of prostate-specific cellular functions requires continuous stimulation by androgens; in prostate cancer tissue, the reciprocal loss of this cellular differentiation, which occurs during progression of the disease, is largely concomitant with a loss of androgen responsiveness by prostatic cells. Identifying the genes involved in either of these largely opposing process, will likely lead to a greater understanding of the fundamental mechanisms involved in both.

Thus far, no genes are known to play a key role in the progressive loss of differentiated phenotype seen in prostate cancer tissue, but various studies indicate the presence of one or more genes on human chromosome 8p that suppress the occurrence and/or progression of the disease. Several investigators have found, based on loss of heterozygosity (LOH) studies, that chromosome bands 8p21 contain loci that are deleted in up to 80% of prostate cancer tissues (Suzuki, *et al.*, *Genes, Chromosomes and Cancer* 13:168-174 (1995), Bova *et al.*, *Cancer Res.* 53:3869-3873 (1993), MacGrogan *et al.*, *Genes, Chromosomes and Cancer* 10:151-159 (1994), Trapman *et al.*, *Cancer Res.* 54:6061-6064 (1994), Macoska, *et al.*, *Cancer Res.* 55:5390-5395 (1995), and Vocke *et al.*, *Cancer Res.* 56:2411-2416 (1996)). In addition, the introduction of human chromosome 8 into the highly metastatic Dunning rat prostate cancer cell line significantly reduces its metastatic potential (Ichikawa *et al.*, *Cancer Res.* 54:2299-2302 (1994)). The loss of 8p during the derivation of subclones from the human prostate cancer line,

LNCaP, is correlated with loss of androgen responsiveness (Konig *et al.*, *Urol. Res.* 17:79-86 (1989)).

Summary of the Invention

5 A prostate-specific human gene (NKX3.1) which maps to 8p21 and encodes a homeodomain-containing protein related to the Drosophila NK gene family was cloned. The gene may play a role in both prostate development and the androgen-driven maintenance of prostatic differentiation in adults. The expression of NKX3.1 in adult humans is restricted to prostate and testes and when assayed
10 in several cell lines, including three lines derived from prostate carcinoma tissue, the gene was expressed solely in the androgen-dependent prostate carcinoma cell line LNCaP. A detailed study of NKX3.1 expression in LNCaP cells has demonstrated that the gene is transcriptionally regulated by androgens. Thus, the new prostate-specific gene NKX3.1 is a candidate for playing a central role in the
15 opposing processes of androgen-driven differentiation of prostatic tissue and loss of that differentiation during the progression of prostate cancer.

 Thus, the present invention provides isolated nucleic acid molecules comprising a polynucleotide encoding the human NK-3 related prostate specific gene 1 (NKX3.1) polypeptide having the amino acid sequence shown in Figure 1
20 or 2 (SEQ ID NOs:2 or 4) or the amino acid sequence encoded by the cDNA clone deposited in a bacterial host as ATCC Deposit Number 209005 on April 28, 1997. The present invention also provides isolated nucleic acid molecules comprising a polynucleotide encoding the human NK-3 related prostate specific gene 1 (HPFCA19) polypeptide having the amino acid sequence shown in Figure
25 5 (SEQ ID NO:9) or the amino acid sequence encoded by the genomic clone deposited in a bacterial host as ATCC Deposit Number 209006 on April 28, 1997.

 The present invention also relates to recombinant vectors, which include the isolated nucleic acid molecules of the present invention, and to host cells

containing the recombinant vectors, as well as to methods of making such vectors and host cells and for using them for production of NKX3.1 polypeptides or peptides by recombinant techniques.

5 The invention further provides an isolated NKX3.1 polypeptide having an amino acid sequence encoded by a polynucleotide described herein.

The present invention also provides a screening method for identifying compounds capable of enhancing or inhibiting a cellular response induced by the NKX3.1, which involves contacting cells which express the NKX3.1 with the candidate compound, assaying a cellular response, and comparing the cellular
10 response to a standard cellular response, the standard being assayed when contact is made in absence of the candidate compound; whereby, an increased cellular response over the standard indicates that the compound is an agonist and a decreased cellular response over the standard indicates that the compound is an antagonist.

15 The invention provides a diagnostic method useful for diagnosis of prostate cancer and other cancers.

An additional aspect of the invention is related to a method for treating an individual in need of an increased level of NKX3.1 activity in the body comprising administering to such an individual a composition comprising a therapeutically
20 effective amount of an isolated NKX3.1 polypeptide of the invention or an agonist thereof.

Brief Description of the Figures

25 Figure 1 shows the nucleotide (SEQ ID NO:1) and deduced amino acid (SEQ ID NO:2) sequences of NKX3.1. Amino acid residues from about 123 to about 153 constitute a homeodomain (underlined region in Figure 1). The protein has a deduced molecular weight of about 26 kDa.

Figure 2 shows the nucleotide (SEQ ID NO:3) and deduced amino acid (SEQ ID NO:4) sequences of NKX3.1. Amino acid residues from about 123 to

about 153 constitute a homeodomain (underlined region in Figure 2). The nucleotide sequence differs from that of SEQ ID NO:1 by one nucleotide. The protein has a deduced molecular weight of about 26 kDa and differs from the protein shown in Figure 1 (SEQ ID NO: 2) by one amino acid.

5 Figure 3 (A) shows the regions of similarity between the amino acid sequences of the NKX3.1 protein NK-3, NK-2 and NK-4 (SEQ ID NOs:5-7). (B) shows the regions of similarity between the amino acid sequences of human and mouse NKX3.1.

10 Figure 4 shows an analysis of the NKX3.1 amino acid sequence. Alpha, beta, turn and coil regions; hydrophilicity and hydrophobicity; amphipathic regions; flexible regions; antigenic index and surface probability are shown. In the "Antigenic Index - Jameson-Wolf" graph, amino acid residues 1-13, 18-24, 35-89, 103-113, 117-130, 173-185 in Figure 1 (SEQ ID NO:2 or 4) correspond to the shown highly antigenic regions of the NKX3.1 protein.

15 Figure 5 shows the nucleotide sequence (SEQ ID NO:8) of the genomic clone of NKX3.1 which include the promoter region. The consensus "CAT" and "TATA" boxes and the first ATG are underlined.

20 Figure 6 shows a schematic representation of the pHE4a expression vector (SEQ ID NO:25). The locations of the kanamycin resistance marker gene, the multiple cloning site linker region, the oriC sequence, and the *lacIq* coding sequence are indicated.

25 Figure 7 shows the nucleotide sequence of the regulatory elements of the pHE4a promoter (SEQ ID NO:26). The two *lac* operator sequences, the Shine-Delgarno sequence (S/D), and the terminal *HindIII* and *NdeI* restriction sites (italicized) are indicated.

Detailed Description

The present invention provides isolated nucleic acid molecules comprising a polynucleotide encoding a NKX3.1 polypeptide having the amino acid sequence

shown in Figures 1 or 2 (SEQ ID NO:2 or SEQ ID NO:4), which was determined by sequencing a cloned cDNA. The NKX3.1 protein of the present invention shares sequence homology with NK-3, NK-2 and NK-4 (Figure 3) (SEQ ID NOs:5-7). The nucleotide sequence shown in Figures 1 or 2 (SEQ ID NO:1 or
5 SEQ ID NO:3) was obtained by sequencing the PSX-lambda-1 (NKX3.1) clone, which was deposited on April 28, 1997 at the American Type Culture Collection, Patent Depository, 10801 University Boulevard, Manassas, VA 20110-2209, and given accession number 209005. The deposited clone is contained in the pBluescript SK(-) plasmid (Stratagene, La Jolla, CA). The nucleotide sequence
10 of the genomic clone shown in Figure 5 (SEQ ID NO:8) was obtained by sequencing the HPFCA19 clone, which was deposited on April 28, 1997 at the American Type Culture Collection Patent Depository, 10801 University Boulevard, Manassas, VA 20110-2209, and given accession number 209006.

Nucleic Acid Molecules

15 Unless otherwise indicated, all nucleotide sequences determined by sequencing a DNA molecule herein were determined using an automated DNA sequencer (such as the Model 373 from Applied Biosystems, Inc.), and all amino acid sequences of polypeptides encoded by DNA molecules determined herein were predicted by translation of a DNA sequence determined as above.
20 Therefore, as is known in the art for any DNA sequence determined by this automated approach, any nucleotide sequence determined herein may contain some errors. Nucleotide sequences determined by automation are typically at least about 90% identical, more typically at least about 95% to at least about 99.9% identical to the actual nucleotide sequence of the sequenced DNA molecule. The
25 actual sequence can be more precisely determined by other approaches including manual DNA sequencing methods well known in the art. As is also known in the art, a single insertion or deletion in a determined nucleotide sequence compared to the actual sequence will cause a frame shift in translation of the nucleotide sequence such that the predicted amino acid sequence encoded by a determined

nucleotide sequence will be completely different from the amino acid sequence actually encoded by the sequenced DNA molecule, beginning at the point of such an insertion or deletion.

Using the information provided herein, such as the nucleotide sequence in
5 Figures 1, 2 or 5, a nucleic acid molecule of the present invention encoding a NKX3.1 polypeptide may be obtained using standard cloning and screening procedures, such as those for cloning cDNAs using mRNA as starting material. Illustrative of the invention, the nucleic acid molecule described in Figures 1, 2 or 5 (SEQ ID NOs:1, 3 or 8) was discovered in a cDNA library derived from normal
10 prostate tissue. The genes were also identified in the following cDNA libraries HO169 (HPFC), S0150 (HPIA). The determined nucleotide sequence of the NKX3.1 cDNAs of Figure 1 or 2 (SEQ ID NOs:1 or 3) contains an open reading frame encoding a protein of 234 amino acid residues, with an initiation codon at positions 1-3 of the nucleotide sequence in Figures 1 or 2 (SEQ ID NOs:1 or 3),
15 and a deduced molecular weight of about 26 kDa. Within the homeodomain, the NKX3.1 protein shown in Figures 1 or 2 (SEQ ID NOs:2 or 4) is about 77 % identical to NK-3, 63% identical to NK-2 and 53% identical to NK-4 (Figure 3A). The nucleotide sequence of the genomic clone is shown in Figure 5 (SEQ ID NO:8).

20 The murine NKX3.1 gene was isolated from a genomic library by hybridization with a human NKX3.1 probe containing the homeobox sequence. Using the human probe to screen a mouse genomic Pst I library, a single strongly-hybridizing colony was identified. Sequence analysis revealed that the human NKX3.1 protein shown in Figure 1 (SEQ ID NO:2) and the mouse homolog
25 shown in Figure 3A are about 43% identical at the N-terminal regions, they have 100% identity in the homeodomain region and are about 67% identical in their C-terminal regions (Figure 3B).

The newly identified gene of the present invention is most closely related to the drosophila NK-3 gene (47/60 aa identity within the homeodomain) and,
30 thus, was named NKX3.1. The longest stretch of complete identity spans aa 40-

60 of the homeodomain and is uniquely conserved between NK-3 and NKX3.1.

This region includes the area between helices II and III of the consensus homeodomain and the entire length of the helix III, suggesting common subtlety in the structure and function of this portion of the drosophila and mammalian NK-3-like genes as compared with the other NK family members. The homeodomain of NKX3.1, is identical to the mouse sequence, and shares block of homology with each of the core Drosophila NK family members, NK-2, NK-3, and NK-4. The longest stretch of homology is within a strongly conserved block from aa 43-58 of the homeodomain including a tyrosine at position 54, which is a common feature of the NK family of genes and their known vertebrate homologues (Lintz, *et al.*, *Development* 119:419-431 (1994); Gehring *et al.*, *Annu. Rev. Biochem.* 63:487-526 (1994)). NKX3.1 and its mouse counterpart represent the first mammalian homologues of the drosophila NK-3 gene.

Typical of most vertebrate homeobox genes, the open reading frame of NKX3.1 is divided between two exons, the second of which contains the entire homeobox region. Each of the analyzed cDNAs terminated approximately 200 Kb beyond the end of the open reading frame and based on the transcription start site predicted from the genomic sequence, this region of the mature mRNA would account for just greater than 1Kb of processed message. However, Northern analyses indicates that the mature NKX3.1 mRNA is approximately 3.5 Kb long (See, Example 4 below) indicating the presence of a long 3' untranslated region (<3.5Kb) which is likely encoded within the second exon, as seen with the newly identified mouse gene.

Thus, the present invention provides a nucleotide sequence encoding the NKX3.1 polypeptide having the amino acid sequence encoded by the cDNA or genomic clone contained in the host identified as ATCC Deposit No. 209005 or 209006 and as shown in Figures 1, 2 or 5 (SEQ ID NOs:2, 4).

As one of ordinary skill would appreciate, due to the possibilities of sequencing errors, the predicted NKX3.1 polypeptide encoded by the deposited

clones comprise about 234 amino acids, but may be anywhere in the range of 220 to 250 amino acids.

As indicated, nucleic acid molecules of the present invention may be in the form of RNA, such as mRNA, or in the form of DNA, including, for instance, cDNA and genomic DNA obtained by cloning or produced synthetically. The DNA may be double-stranded or single-stranded. Single-stranded DNA or RNA may be the coding strand, also known as the sense strand, or it may be the non-coding strand, also referred to as the anti-sense strand.

By "isolated" nucleic acid molecule(s) is intended a nucleic acid molecule, DNA or RNA, which has been removed from its native environment. For example, recombinant DNA molecules contained in a vector are considered isolated for the purposes of the present invention. Further examples of isolated DNA molecules include recombinant DNA molecules maintained in heterologous host cells or purified (partially or substantially) DNA molecules in solution. Isolated RNA molecules include *in vivo* or *in vitro* RNA transcripts of the DNA molecules of the present invention. Isolated nucleic acid molecules according to the present invention further include such molecules produced synthetically.

Isolated nucleic acid molecules of the present invention include DNA molecules comprising an open reading frame (ORF) shown in Figures 1, 2 or 5 (SEQ ID NOs:1, 3 or 8); DNA molecules comprising the coding sequence for the NKX3.1 protein shown in Figures 1, 2 (SEQ ID NOs:2 or 4); and DNA molecules which comprise a sequence substantially different from those described above but which, due to the degeneracy of the genetic code, still encode the NKX3.1 protein. Of course, the genetic code is well known in the art. Thus, it would be routine for one skilled in the art to generate such degenerate variants.

In addition, the invention provides nucleic acid molecules having nucleotide sequences related to extensive portions of SEQ ID NO:1 or 3 which have been determined from the following related Genbank cDNA clones: AA492170 (SEQ ID NO:27) and AA855030 (SEQ ID NO:28).

In addition, the invention provides nucleic acid molecules having nucleotide sequences related to extensive portions of SEQ ID NO:1 or 3 which have been determined from the following related cDNA clones: HPIAA91R (SEQ ID NO:29) and HAWAU13R (SEQ ID NO:30).

5 In another aspect, the invention provides isolated nucleic acid molecules encoding the NKX3.1 polypeptide having an amino acid sequence encoded by the deposited clones contained in the plasmid deposited as ATCC Deposit No. 209005 or 209006 on April 28, 1997. Preferably, this nucleic acid molecule will encode the polypeptide encoded by the above-described deposited clones. The
10 invention further provides an isolated nucleic acid molecule having the nucleotide sequence shown in Figures 1, 2 or 5 (SEQ ID NOs:1, 3 or 8) or the nucleotide sequence of the NKX3.1 cDNA or genomic sequence contained in the above-described deposited clones, or a nucleic acid molecule having a sequence complementary to one of the above sequences. Such isolated molecules,
15 particularly DNA molecules, are useful as probes for gene mapping, by *in situ* hybridization with chromosomes, and for detecting expression of the NKX3.1 gene in human tissue, for instance, by Northern blot analysis.

The present invention is further directed to fragments of the isolated nucleic acid molecules described herein. By a fragment of an isolated nucleic acid
20 molecule having the nucleotide sequence of the deposited clones or the nucleotide sequence shown in Figures 1, 2 or 5 (SEQ ID NOs:1, 3 or 8) is intended fragments at least about 15 nt, and more preferably at least about 20 nt, still more preferably at least about 30 nt, and even more preferably, at least about 40 nt in length which are useful as diagnostic probes and primers as discussed herein. Of
25 course, larger fragments 50, 75, 100, 125, 150, 175, 200, 225, 250, 300, 325, 350, 375, 400, 425, 450, 475, 500, 525, 550, 575, 600, 625, 650, 675, or 700 nt in length are also useful according to the present invention as are fragments corresponding to most, if not all, of the nucleotide sequence of the deposited clones or as shown in Figures 1, 2 or 5 (SEQ ID NOs:1, 3 or 8). By a fragment
30 at least 20 nt in length, for example, is intended fragments which include 20 or

more contiguous bases from the nucleotide sequence of the deposited clones or the nucleotide sequence as shown in Figures 1, 2 or 5 (SEQ ID NOs:1, 3 or 8).

Preferred nucleic acid fragments of the present invention include nucleic acid molecules encoding epitope-bearing portions of the NKX3.1 protein. In particular, such nucleic acid fragments of the present invention include nucleic acid molecules encoding: a polypeptide comprising amino acid residues from about 1 to about 13 in Figures 1 or 2 (SEQ ID NOs:2 or 4); a polypeptide comprising amino acid residues from about 18 to about 24 in Figures 1 or 2 (SEQ ID NOs:2 or 4); a polypeptide comprising amino acid residues from about 35 to about 89 in Figures 1 or 2 (SEQ ID NOs:2 or 4); a polypeptide comprising amino acid residues from about 103 to about 113 in Figures 1 or 2 (SEQ ID NOs:2 or 4); a polypeptide comprising amino acid residues from about 117 to about 130 in Figures 1 or 2 (SEQ ID NOs:2 or 4); and a polypeptide comprising amino acid residues from about 173 to about 185 in Figures 1 or 2 (SEQ ID NO:2 or 4). The inventors have determined that the above polypeptide fragments are antigenic regions of the NKX3.1 protein. Methods for determining other such epitope-bearing portions of the NKX3.1 protein are described in detail below.

In another aspect, the invention provides an isolated nucleic acid molecule comprising a polynucleotide which hybridizes under stringent hybridization conditions to a portion of the polynucleotide in a nucleic acid molecule of the invention described above, for instance, the deposited clones contained in ATCC Deposit 209005 or 209006. By "stringent hybridization conditions" is intended overnight incubation at 42°C in a solution comprising: 50% formamide, 5x SSC (150 mM NaCl, 15mM trisodium citrate), 50 mM sodium phosphate (pH 7.6), 5x Denhardt's solution, 10% dextran sulfate, and 20 g/ml denatured, sheared salmon sperm DNA, followed by washing the filters in 0.1x SSC at about 65°C.

By a polynucleotide which hybridizes to a "portion" of a polynucleotide is intended a polynucleotide (either DNA or RNA) hybridizing to at least about 15 nucleotides (nt), and more preferably at least about 20 nt, still more preferably at least about 30 nt, and even more preferably about 30-70 nt of the reference

polynucleotide. These are useful as diagnostic probes and primers as discussed above and in more detail below.

By a portion of a polynucleotide of "at least 20 nt in length," for example, is intended 20 or more contiguous nucleotides from the nucleotide sequence of the reference polynucleotide (e.g., the deposited clones or the nucleotide sequence as shown in Figures 1, 2 or 5 (SEQ ID NOs: 1, 3 or 8). Of course, a polynucleotide which hybridizes only to a poly A sequence (such as the 3' terminal poly(A) tract of the NKX3.1 cDNA shown in Figures 1, 2 or 5 (SEQ ID NOs: 1, 3 or 8), or to a complementary stretch of T (or U) residues, would not be included in a polynucleotide of the invention used to hybridize to a portion of a nucleic acid of the invention, since such a polynucleotide would hybridize to any nucleic acid molecule containing a poly (A) stretch or the complement thereof (e.g., practically any double-stranded cDNA clone).

As indicated, nucleic acid molecules of the present invention which encode a NKX3.1 polypeptide may include, but are not limited to those encoding the amino acid sequence of the polypeptide, by itself; the coding sequence for the polypeptide and additional sequences, such as those encoding an amino acid leader or secretory sequence, such as a pre-, or pro- or prepro- protein sequence; the coding sequence of the polypeptide, with or without the aforementioned additional coding sequences, together with additional, non-coding sequences, including for example, but not limited to introns and non-coding 5' and 3' sequences, such as the transcribed, non-translated sequences that play a role in transcription, mRNA processing, including splicing and polyadenylation signals, for example - ribosome binding and stability of mRNA; an additional coding sequence which codes for additional amino acids, such as those which provide additional functionalities. Thus, the sequence encoding the polypeptide may be fused to a marker sequence, such as a sequence encoding a peptide which facilitates purification of the fused polypeptide. In certain preferred embodiments of this aspect of the invention, the marker amino acid sequence is a hexa-histidine peptide, such as the tag provided in a pQE vector (Qiagen, Inc.), among others, many of which are commercially

available. As described in Gentz *et al.*, *Proc. Natl. Acad. Sci. USA* 86:821-824 (1989), for instance, hexa-histidine provides for convenient purification of the fusion protein. The "HA" tag is another peptide useful for purification which corresponds to an epitope derived from the influenza hemagglutinin protein, which has been described by Wilson *et al.*, *Cell* 37: 767 (1984). As discussed below, other such fusion proteins include the NKX3.1 fused to Fc at the N- or C-terminus.

The present invention further relates to variants of the nucleic acid molecules of the present invention, which encode portions, analogs or derivatives of the NKX3.1 protein. Variants may occur naturally, such as a natural allelic variant. By an "allelic variant" is intended one of several alternate forms of a gene occupying a given locus on a chromosome of an organism. *Genes II*, Lewin, B., ed., John Wiley & Sons, New York (1985). Non-naturally occurring variants may be produced using art-known mutagenesis techniques.

Such variants include those produced by nucleotide substitutions, deletions or additions, which may involve one or more nucleotides. The variants may be altered in coding regions, non-coding regions, or both. Alterations in the coding regions may produce conservative or non-conservative amino acid substitutions, deletions or additions. Especially preferred among these are silent substitutions, additions and deletions, which do not alter the properties and activities of the NKX3.1 protein or portions thereof. Also especially preferred in this regard are conservative substitutions.

Further embodiments of the invention include isolated nucleic acid molecules comprising a polynucleotide having a nucleotide sequence at least 95%, 96%, 97%, 98% or 99% identical to (a) a nucleotide sequence encoding the polypeptide having the amino acid sequence in SEQ ID NOs:2 or 4; (b) a nucleotide sequence encoding the polypeptide having the amino acid sequence in SEQ ID NOs:2 or 4, but lacking the N-terminal methionine; (c) a nucleotide sequence encoding the full-length NKX3.1 polypeptide having the complete amino acid sequence encoded by the clone contained in ATCC Deposit No. 209005 or

209006; (d) a nucleotide sequence encoding the homeodomain of the NKX3.1 polypeptide (predicted to constitute amino acid residues 123-153 in Figures 1 or 2 [SEQ ID NOs: 2 or 4]); or (e) a nucleotide sequence complementary to any of the nucleotide sequences in (a), (b), (c) or (d).

5 By a polynucleotide having a nucleotide sequence at least, for example, 95% "identical" to a reference nucleotide sequence encoding a NKX3.1 polypeptide is intended that the nucleotide sequence of the polynucleotide is identical to the reference sequence except that the polynucleotide sequence may include up to five point mutations per each 100 nucleotides of the reference
10 nucleotide sequence encoding the NKX3.1 polypeptide. In other words, to obtain a polynucleotide having a nucleotide sequence at least 95% identical to a reference nucleotide sequence, up to 5% of the nucleotides in the reference sequence may be deleted or substituted with another nucleotide, or a number of nucleotides up to 5% of the total nucleotides in the reference sequence may be inserted into the
15 reference sequence. These mutations of the reference sequence may occur at the 5' or 3' terminal positions of the reference nucleotide sequence or anywhere between those terminal positions, interspersed either individually among nucleotides in the reference sequence or in one or more contiguous groups within the reference sequence.

20 As a practical matter, whether any particular nucleic acid molecule is at least 95%, 96%, 97%, 98% or 99% identical to, for instance, the nucleotide sequence shown in Figures 1, 2 or 5 or to the nucleotides sequence of the deposited clones can be determined conventionally using known computer programs such as the Bestfit program (Wisconsin Sequence Analysis Package,
25 Version 8 for Unix, Genetics Computer Group, University Research Park, 575 Science Drive, Madison, WI 53711. Bestfit uses the local homology algorithm of Smith and Waterman, *Advances in Applied Mathematics* 2: 482-489 (1981), to find the best segment of homology between two sequences. When using Bestfit or any other sequence alignment program to determine whether a particular
30 sequence is, for instance, 95% identical to a reference sequence according to the

present invention, the parameters are set, of course, such that the percentage of identity is calculated over the full length of the reference nucleotide sequence and that gaps in homology of up to 5% of the total number of nucleotides in the reference sequence are allowed.

5 The present application is directed to nucleic acid molecules at least 95%, 96%, 97%, 98% or 99% identical to the nucleic acid sequence shown in Figures 1, 2 or 5 (SEQ ID NOs:1, 3 or 8) or to the nucleic acid sequence of the deposited clones, irrespective of whether they encode a polypeptide having NKX3.1 activity. This is because even where a particular nucleic acid molecule does not
10 encode a polypeptide having NKX3.1 activity, one of skill in the art would still know how to use the nucleic acid molecule, for instance, as a hybridization probe or a polymerase chain reaction (PCR) primer. Uses of the nucleic acid molecules of the present invention that do not encode a polypeptide having NKX3.1 activity include, *inter alia*, (1) isolating the NKX3.1 gene or allelic variants thereof in a
15 cDNA library; (2) *in situ* hybridization (e.g., "FISH") to metaphase chromosomal spreads to provide precise chromosomal location of the NKX3.1 gene, as described in Verma *et al.*, *Human Chromosomes: A Manual of Basic Techniques*, Pergamon Press, New York (1988); and Northern Blot analysis for detecting NKX3.1 mRNA expression in specific tissues.

20 Preferred, however, are nucleic acid molecules having sequences at least 95%, 96%, 97%, 98% or 99% identical to the nucleic acid sequence shown in Figures 1, 2 or 5 (SEQ ID NOs:1, 3 or 8) or to the nucleic acid sequence of the deposited clones which do, in fact, encode a polypeptide having NKX3.1 protein activity. By "a polypeptide having NKX3.1 activity" is intended polypeptides
25 exhibiting activity similar, but not necessarily identical, to an activity of the NKX3.1 protein of the invention, as measured in a particular biological assay. For example, NKX3.1 protein activity can be measured using the DNA binding experiment set forth in Example 7 below.

30 Of course, due to the degeneracy of the genetic code, one of ordinary skill in the art will immediately recognize that a large number of the nucleic acid

5 molecules having a sequence at least 95%, 96%, 97%, 98%, or 99% identical to the nucleic acid sequence of the deposited clones or the nucleic acid sequence shown in Figures 1, 2 or 5 (SEQ ID NOs:1, 3 or 8) will encode a polypeptide "having NKX3.1 protein activity." In fact, since degenerate variants of these nucleotide sequences all encode the same polypeptide, this will be clear to the skilled artisan even without performing the above described comparison assay. It will be further recognized in the art that, for such nucleic acid molecules that are not degenerate variants, a reasonable number will also encode a polypeptide having NKX3.1 protein activity. This is because the skilled artisan is fully aware of amino acid substitutions that are either less likely or not likely to significantly effect protein function (e.g., replacing one aliphatic amino acid with a second aliphatic amino acid).

10 For example, guidance concerning how to make phenotypically silent amino acid substitutions is provided in Bowie, J. U. *et al.*, "Deciphering the Message in Protein Sequences: Tolerance to Amino Acid Substitutions," *Science* 247:1306-1310 (1990), wherein the authors indicate that proteins are surprisingly tolerant of amino acid substitutions.

Vectors and Host Cells

20 The present invention also relates to vectors which include the isolated DNA molecules of the present invention, host cells which are genetically engineered with the recombinant vectors, and the production of NKX3.1 polypeptides or fragments thereof by recombinant techniques.

25 The polynucleotides may be joined to a vector containing a selectable marker for propagation in a host. Generally, a plasmid vector is introduced in a precipitate, such as a calcium phosphate precipitate, or in a complex with a charged lipid. If the vector is a virus, it may be packaged *in vitro* using an appropriate packaging cell line and then transduced into host cells.

The DNA insert should be operatively linked to an appropriate promoter, such as the phage lambda PL promoter, the *E. coli lac*, *trp* and *tac* promoters, the SV40 early and late promoters and promoters of retroviral LTRs, to name a few. Other suitable promoters will be known to the skilled artisan. The expression constructs will further contain sites for transcription initiation, termination and, in the transcribed region, a ribosome binding site for translation. The coding portion of the mature transcripts expressed by the constructs will preferably include a translation initiating at the beginning and a termination codon (UAA, UGA or UAG) appropriately positioned at the end of the polypeptide to be translated.

As indicated, the expression vectors will preferably include at least one selectable marker. Such markers include dihydrofolate reductase or neomycin resistance for eukaryotic cell culture and tetracycline or ampicillin resistance genes for culturing in *E. coli* and other bacteria. Representative examples of appropriate hosts include, but are not limited to, bacterial cells, such as *E. coli*, *Streptomyces* and *Salmonella typhimurium* cells; fungal cells, such as yeast cells; insect cells such as *Drosophila* S2 and *Spodoptera* Sf9 cells; animal cells such as CHO, COS and Bowes melanoma cells; and plant cells. Appropriate culture mediums and conditions for the above-described host cells are known in the art.

In addition to the use of expression vectors in the practice of the present invention, the present invention further includes novel expression vectors comprising operator and promoter elements operatively linked to nucleotide sequences encoding a protein of interest. One example of such a vector is pHE4a which is described in detail below.

As summarized in Figures 6 and 7, components of the pHE4a vector (SEQ ID NO:25) include: 1) a neomycinphosphotransferase gene as a selection marker, 2) an *E. coli* origin of replication, 3) a T5 phage promoter sequence, 4) two *lac* operator sequences, 5) a Shine-Delgarno sequence, 6) the lactose operon repressor gene (*lacIq*) and 7) a multiple cloning site linker region. The origin of replication (*oriC*) is derived from pUC19 (LTI, Gaithersburg, MD). The promoter sequence and operator sequences were made synthetically. Synthetic production

of nucleic acid sequences is well known in the art. CLONTECH 95/96 Catalog, pages 215-216, CLONTECH, 1020 East Meadow Circle, Palo Alto, CA 94303. The pHE4a vector was deposited with the ATCC on February 25, 1998, and given accession number 209645.

5 A nucleotide sequence encoding NKX3.1 (SEQ ID Nos: 1 or 3), is operatively linked to the promoter and operator of pHE4a by restricting the vector with NdeI and either XbaI, BamHI, XhoI, or Asp718, and isolating the larger fragment (the multiple cloning site region is about 310 nucleotides) on a gel. The nucleotide sequence encoding NKX3.1 (SEQ ID NO:1 or 3) having the
10 appropriate restriction sites is generated, for example, according to the PCR protocol described in Example 1, using PCR primers having restriction sites for NdeI (as the 5' primer) and either XbaI, BamHI, XhoI, or Asp718 (as the 3' primer). The PCR insert is gel purified and restricted with compatible enzymes. The insert and vector are ligated according to standard protocols.

15 As noted above, the pHE4a vector contains a *lacIq* gene. *LacIq* is an allele of the *lacI* gene which confers tight regulation of the *lac* operator. Amann, E. *et al.*, *Gene* 69:301-315 (1988); Stark, M., *Gene* 51:255-267 (1987). The *lacIq* gene encodes a repressor protein which binds to *lac* operator sequences and blocks transcription of down-stream (*i.e.*, 3') sequences. However, the *lacIq* gene
20 product dissociates from the *lac* operator in the presence of either lactose or certain lactose analogs, *e.g.*, isopropyl B-D-thiogalactopyranoside (IPTG). NKX3.1 thus is not produced in appreciable quantities in uninduced host cells containing the pHE4a vector. Induction of these host cells by the addition of an agent such as IPTG, however, results in the expression of the NKX3.1 coding
25 sequence.

 The promoter/operator sequences of the pHE4a vector (SEQ ID NO:26) comprise a T5 phage promoter and two *lac* operator sequences. One operator is located 5' to the transcriptional start site and the other is located 3' to the same site. These operators, when present in combination with the *lacIq* gene product,
30 confer tight repression of down-stream sequences in the absence of a *lac* operon

inducer, *e.g.*, IPTG. Expression of operatively linked sequences located down-stream from the *lac* operators may be induced by the addition of a *lac* operon inducer, such as IPTG. Binding of a *lac* inducer to the *lacI* proteins results in their release from the *lac* operator sequences and the initiation of transcription of operatively linked sequences. *Lac* operon regulation of gene expression is reviewed in Devlin, T., TEXTBOOK OF BIOCHEMISTRY WITH CLINICAL CORRELATIONS, 4th Edition (1997), pages 802-807.

The pHE4 series of vectors contain all of the components of the pHE4a vector except for the NKX3.1 coding sequence. Features of the pHE4a vectors include optimized synthetic T5 phage promoter, *lac* operator, and Shine-Delgarno sequences. Further, these sequences are also optimally spaced so that expression of an inserted gene may be tightly regulated and high level of expression occurs upon induction.

Among known bacterial promoters suitable for use in the production of proteins of the present invention include the *E. coli lacI* and *lacZ* promoters, the T3 and T7 promoters, the *gpt* promoter, the lambda PR and PL promoters and the *trp* promoter. Suitable eukaryotic promoters include the CMV immediate early promoter, the HSV thymidine kinase promoter, the early and late SV40 promoters, the promoters of retroviral LTRs, such as those of the Rous Sarcoma Virus (RSV), and metallothionein promoters, such as the mouse metallothionein-I promoter.

The pHE4a vector also contains a Shine-Delgarno sequence 5' to the AUG initiation codon. Shine-Delgarno sequences are short sequences generally located about 10 nucleotides up-stream (*i.e.*, 5') from the AUG initiation codon. These sequences essentially direct prokaryotic ribosomes to the AUG initiation codon.

Thus, the present invention is also directed to expression vector useful for the production of the proteins of the present invention. This aspect of the invention is exemplified by the pHE4a vector (SEQ ID NO:25).

Among vectors preferred for use in bacteria include pQE70, pQE60 and pQE-9, available from Qiagen; pBS vectors, Phagescript vectors, Bluescript

vectors, pNH8A, pNH16a, pNH18A, pNH46A, available from Stratagene; and ptrc99a, pKK223-3, pKK233-3, pDR540, pRIT5 available from Pharmacia. Among preferred eukaryotic vectors are pWLNEO, pSV2CAT, pOG44, pXT1 and pSG available from Stratagene; and pSVK3, pBPV, pMSG and pSVL available from Pharmacia. Other suitable vectors will be readily apparent to the skilled artisan.

Introduction of the construct into the host cell can be effected by calcium phosphate transfection, DEAE-dextran mediated transfection, cationic lipid-mediated transfection, electroporation, transduction, infection or other methods. Such methods are described in many standard laboratory manuals, such as Davis *et al.*, *Basic Methods In Molecular Biology* (1986).

The polypeptide may be expressed in a modified form, such as a fusion protein, and may include not only secretion signals, but also additional heterologous functional regions. For instance, a region of additional amino acids, particularly charged amino acids, may be added to the N-terminus of the polypeptide to improve stability and persistence in the host cell, during purification, or during subsequent handling and storage. Also, peptide moieties may be added to the polypeptide to facilitate purification. Such regions may be removed prior to final preparation of the polypeptide. The addition of peptide moieties to polypeptides to engender secretion or excretion, to improve stability and to facilitate purification, among others, are familiar and routine techniques in the art. A preferred fusion protein comprises a heterologous region from immunoglobulin that is useful to solubilize proteins. For example, EP-A-O 464 533 (Canadian counterpart 2045869) discloses fusion proteins comprising various portions of constant region of immunoglobulin molecules together with another human protein or part thereof. In many cases, the Fc part in a fusion protein is thoroughly advantageous for use in therapy and diagnosis and thus results, for example, in improved pharmacokinetic properties (EP-A 0232 262). On the other hand, for some uses it would be desirable to be able to delete the Fc part after the fusion protein has been expressed, detected and purified in the advantageous

manner described. This is the case when Fc portion proves to be a hindrance to use in therapy and diagnosis, for example when the fusion protein is to be used as antigen for immunizations. In drug discovery, for example, human proteins, such as, hIL5-receptor has been fused with Fc portions for the purpose of high-throughput screening assays to identify antagonists of hIL-5. See, D. Bennett *et al.*, *Journal of Molecular Recognition*, Vol. 8:52-58 (1995) and K. Johanson *et al.*, *The Journal of Biological Chemistry*, Vol. 270, No. 16:9459-9471 (1995).

The NKX3.1 protein can be recovered and purified from recombinant cell cultures by well-known methods including ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite chromatography and lectin chromatography. Most preferably, high performance liquid chromatography ("HPLC") is employed for purification. Polypeptides of the present invention include naturally purified products, products of chemical synthetic procedures, and products produced by recombinant techniques from a prokaryotic or eukaryotic host, including, for example, bacterial, yeast, higher plant, insect and mammalian cells. Depending upon the host employed in a recombinant production procedure, the polypeptides of the present invention may be glycosylated or may be non-glycosylated. In addition, polypeptides of the invention may also include an initial modified methionine residue, in some cases as a result of host-mediated processes.

NKX3.1 Polypeptides and Fragments

The invention further provides an isolated NKX3.1 polypeptide having the amino acid sequence encoded by the deposited clones, or the amino acid sequence in Figures 1 or 2 (SEQ ID NOs:2 or 4), or a peptide or polypeptide comprising a portion of the above polypeptides.

It will be recognized in the art that some amino acid sequences of the NKX3.1 polypeptide can be varied without significant effect of the structure or function of the protein. If such differences in sequence are contemplated, it should be remembered that there will be critical areas on the protein which determine activity.

Thus, the invention further includes variations of the NKX3.1 polypeptide which show substantial NKX3.1 polypeptide activity or which include regions of NKX3.1 protein such as the protein portions discussed below. Such mutants include deletions, insertions, inversions, repeats, and type substitutions. As indicated above, guidance concerning which amino acid changes are likely to be phenotypically silent can be found in Bowie, J.U., *et al.*, "Deciphering the Message in Protein Sequences: Tolerance to Amino Acid Substitutions," *Science* 247:1306-1310 (1990).

Thus, the fragment, derivative or analog of the polypeptide of Figures 1 or 2 (SEQ ID NOs:2 or 4), or that encoded by the deposited cDNA or genomic clones, may be (i) one in which one or more of the amino acid residues are substituted with a conserved or non-conserved amino acid residue (preferably a conserved amino acid residue) and such substituted amino acid residue may or may not be one encoded by the genetic code, or (ii) one in which one or more of the amino acid residues includes a substituent group, or (iii) one in which the polypeptide is fused with another compound, such as a compound to increase the half-life of the polypeptide (for example, polyethylene glycol), or (iv) one in which the additional amino acids are fused to the mature polypeptide, such as an IgG Fc fusion region peptide or leader or secretory sequence or a sequence which is employed for purification of the polypeptide or a proprotein sequence. Such fragments, derivatives and analogs are deemed to be within the scope of those skilled in the art from the teachings herein.

Of particular interest are substitutions of charged amino acids with another charged amino acid and with neutral or negatively charged amino acids. The latter results in proteins with reduced positive charge to improve the characteristics of

the NKX3.1 protein. The prevention of aggregation is highly desirable. Aggregation of proteins not only results in a loss of activity but can also be problematic when preparing pharmaceutical formulations, because they can be immunogenic. (Pinckard *et al.*, *Clin Exp. Immunol.* 2:331-340 (1967); Robbins *et al.*, *Diabetes* 36:838-845 (1987); Cleland *et al.* *Crit. Rev. Therapeutic Drug Carrier Systems* 10:307-377 (1993)).

As indicated, changes are preferably of a minor nature, such as conservative amino acid substitutions that do not significantly affect the folding or activity of the protein (see Table 1).

TABLE 1. Conservative Amino Acid Substitutions.

Aromatic	Phenylalanine Tryptophan Tyrosine
Hydrophobic	Leucine Isoleucine Valine
Polar	Glutamine Asparagine
Basic	Arginine Lysine Histidine
Acidic	Aspartic Acid Glutamic Acid
Small	Alanine Serine Threonine Methionine Glycine

Of course, the number of amino acid substitutions a skilled artisan would make depends on many factors, including those described above. Generally speaking, the number of amino acid substitutions for any given NKX3.1 polypeptide will not be more than 50, 40, 30, 20, 10, 5, or 3.

Amino acids in the NKX3.1 protein of the present invention that are essential for function can be identified by methods known in the art, such as site-directed mutagenesis or alanine-scanning mutagenesis (Cunningham and Wells, *Science* 244:1081-1085 (1989)). The latter procedure introduces single alanine mutations at every residue in the molecule. The resulting mutant molecules are then tested for biological activity such as receptor binding or *in vitro*, or *in vitro* proliferative activity. Sites that are critical for NKX3.1-DNA binding can also be determined by structural analysis such as crystallization, nuclear magnetic resonance or photoaffinity labeling (Smith *et al.*, *J. Mol. Biol.* 224:899-904 (1992) and de Vos *et al.* *Science* 255:306-312 (1992)).

The polypeptides of the present invention are preferably provided in an isolated form, and preferably are substantially purified. By "isolated polypeptide" is intended a polypeptide removed from its native environment. Thus, a polypeptide produced and/or contained within a recombinant host cell is considered isolated for purposes of the present invention. Also intended as an "isolated polypeptide" are polypeptides that have been purified, partially or substantially, from a recombinant host cell. For example, a recombinantly produced version of the NKX3.1 polypeptide can be substantially purified by the one-step method described in Smith and Johnson, *Gene* 67:31-40 (1988).

The polypeptides of the present invention include the polypeptide encoded by the deposited cDNA including the leader; a polypeptide comprising amino acids about 1 to about 234 in SEQ ID NOs:2 or 4; a polypeptide comprising amino acids about 2 to about 234 in SEQ ID NOs:2 or 4; a polypeptide comprising amino acids about 2 to about 365 in SEQ ID NOs:2 or 4 but lacking the homeodomain of the polypeptide of SEQ ID NOs:2 or 4 as well as polypeptides which are at least 95% identical, still more preferably at least 96%, 97%, 98% or 99% identical to the polypeptides encoded by the deposited clones, to the polypeptide of Figures 1 or 2 (SEQ ID NOs:2 or 4), and also include portions of such polypeptides with at least 30 amino acids and more preferably at least 50 amino acids.

By a polypeptide having an amino acid sequence at least, for example, 95% "identical" to a reference amino acid sequence of a NKX3.1 polypeptide is intended that the amino acid sequence of the polypeptide is identical to the reference sequence except that the polypeptide sequence may include up to five amino acid alterations per each 100 amino acids of the reference amino acid of the NKX3.1 polypeptide. In other words, to obtain a polypeptide having an amino acid sequence at least 95% identical to a reference amino acid sequence, up to 5% of the amino acid residues in the reference sequence may be deleted or substituted with another amino acid, or a number of amino acids up to 5% of the total amino acid residues in the reference sequence may be inserted into the reference sequence. These alterations of the reference sequence may occur at the amino or carboxy terminal positions of the reference amino acid sequence or anywhere between those terminal positions, interspersed either individually among residues in the reference sequence or in one or more contiguous groups within the reference sequence.

As a practical matter, whether any particular polypeptide is at least 95%, 96%, 97%, 98% or 99% identical to, for instance, the amino acid sequence shown in Figures 1 or 2 (SEQ ID NOs:2 or 4) or to the amino acid sequence encoded by deposited clones can be determined conventionally using known computer programs such the Bestfit program (Wisconsin Sequence Analysis Package, Version 8 for Unix, Genetics Computer Group, University Research Park, 575 Science Drive, Madison, WI 53711. When using Bestfit or any other sequence alignment program to determine whether a particular sequence is, for instance, 95% identical to a reference sequence according to the present invention, the parameters are set, of course, such that the percentage of identity is calculated over the full length of the reference amino acid sequence and that gaps in homology of up to 5% of the total number of amino acid residues in the reference sequence are allowed.

The polypeptide of the present invention could be used as a molecular weight marker on SDS-PAGE gels or on molecular sieve gel filtration columns using methods well known to those of skill in the art.

5 In another aspect, the invention provides a peptide or polypeptide comprising an epitope-bearing portion of a polypeptide of the invention. The epitope of this polypeptide portion is an immunogenic or antigenic epitope of a polypeptide described herein. An "immunogenic epitope" is defined as a part of a protein that elicits an antibody response when the whole protein is the immunogen. On the other hand, a region of a protein molecule to which an
10 antibody can bind is defined as an "antigenic epitope." The number of immunogenic epitopes of a protein generally is less than the number of antigenic epitopes. See, for instance, Geysen *et al.*, *Proc. Natl. Acad. Sci. USA* 81:3998-4002 (1983).

As to the selection of peptides or polypeptides bearing an antigenic epitope
15 (i.e., that contain a region of a protein molecule to which an antibody can bind), it is well known in that art that relatively short synthetic peptides that mimic part of a protein sequence are routinely capable of eliciting an antiserum that reacts with the partially mimicked protein. See, for instance, Sutcliffe, J. G., Shinnick, T. M., Green, N. and Learner, R.A. (1983) Antibodies that react with
20 predetermined sites on proteins. *Science* 219:660-666. Peptides capable of eliciting protein-reactive sera are frequently represented in the primary sequence of a protein, can be characterized by a set of simple chemical rules, and are confined neither to immunodominant regions of intact proteins (i.e., immunogenic epitopes) nor to the amino or carboxyl terminals.

25 Antigenic epitope-bearing peptides and polypeptides of the invention are therefore useful to raise antibodies, including monoclonal antibodies, that bind specifically to a polypeptide of the invention. See, for instance, Wilson *et al.*, *Cell* 37:767-778 (1984) at 777.

30 Antigenic epitope-bearing peptides and polypeptides of the invention preferably contain a sequence of at least seven, more preferably at least nine and

most preferably between about at least about 15 to about 30 amino acids contained within the amino acid sequence of a polypeptide of the invention.

Non-limiting examples of antigenic polypeptides or peptides that can be used to generate NKX3.1-specific antibodies include: a polypeptide comprising amino acid residues from about 1 to about 13 in SEQ ID NO:2 or SEQ ID NO:4; a polypeptide comprising amino acid residues from about 18 to about 24 in SEQ ID NO:2 or SEQ ID NO:4; a polypeptide comprising amino acid residues from about 35 to about 89 in SEQ ID NO:2 or SEQ ID NO:4; a polypeptide comprising amino acid residues from about 103 to about 113 in SEQ ID NO:2 or SEQ ID NO:4; a polypeptide comprising amino acid residues from about 117 to about 130 in SEQ ID NO:2 or SEQ ID NO:4; and a polypeptide comprising amino acid residues from about 173 to about 185 in SEQ ID NO:2 or SEQ ID NO:4. As indicated above, the inventors have determined that the above polypeptide fragments are antigenic regions of the NKX3.1 protein.

The epitope-bearing peptides and polypeptides of the invention may be produced by any conventional means. Houghten, R. A. (1985) General method for the rapid solid-phase synthesis of large numbers of peptides: specificity of antigen-antibody interaction at the level of individual amino acids. *Proc. Natl. Acad. Sci. USA* 82:5131-5135. This "Simultaneous Multiple Peptide Synthesis (SMPS)" process is further described in U.S. Patent No. 4,631,211 to Houghten *et al.* (1986).

As one of skill in the art will appreciate, NKX3.1 polypeptides of the present invention and the epitope-bearing fragments thereof described above can be combined with parts of the constant domain of immunoglobulins (IgG), resulting in chimeric polypeptides. These fusion proteins facilitate purification and show an increased half-life *in vivo*. This has been shown, e.g., for chimeric proteins consisting of the first two domains of the human CD4-polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins (EPA 394,827; Traunecker *et al.*, *Nature* 331:84- 86 (1988)). Fusion proteins that have a disulfide-linked dimeric structure due to the IgG part

can also be more efficient in binding and neutralizing other molecules than the monomeric NKX3.1 protein or protein fragment alone (Fountoulakis *et al.*, *J. Biochem* 270:3958-3964 (1995)).

N-terminal and C-terminal Deletion Mutants

5 In one embodiment, the present invention provides polypeptides having one or more residues deleted from the amino terminus of the amino acid sequence of the NKX3 polypeptide depicted in Figure 1, Figure 2, or encoded by the cDNA of the deposited clone. Particularly, in one embodiment, N-terminal deletions of the NKX3 polypeptide can be described by the general formula m to 234, where
10 m is any one of the integers from 2 to 233 corresponding to the position of the amino acid residue identified in SEQ ID NO:2 and, preferably, corresponds to one of the N-terminal amino acid residues identified in the N-terminal deletions specified herein. In specific embodiments, N-terminal deletions of the NKX3.1 polypeptide of the invention comprise, or alternatively consist of, amino acid
15 residues: L-2 to G-234; R-3 to G-234; V-4 to G-234; P-5 to G-234; E-6 to G-234; P-7 to G-234; R-8 to G-234; P-9 to G-234; G-10 to G-234; E-11 to G-234; A-12 to G-234; K-13 to G-234; A-14 to G-234; E-15 to G-234; G-16 to G-234; A-17 to G-234; A-18 to G-234; P-19 to G-234; P-20 to G-234; T-21 to G-234; P-22 to G-234; S-23 to G-234; K-24 to G-234; P-25 to G-234; L-26 to
20 G-234; T-27 to G-234; S-28 to G-234; F-29 to G-234; L-30 to G-234; I-31 to G-234; Q-32 to G-234; D-33 to G-234; I-34 to G-234; L-35 to G-234; R-36 to G-234; D-37 to G-234; G-38 to G-234; A-39 to G-234; Q-40 to G-234; R-41 to G-234; Q-42 to G-234; G-43 to G-234; G-44 to G-234; R-45 to G-234; T-46 to G-234; S-47 to G-234; S-48 to G-234; Q-49 to G-234; R-50 to G-234; Q-51 to
25 G-234; R-52 to G-234; D-53 to G-234; P-54 to G-234; E-55 to G-234; P-56 to G-234; E-57 to G-234; P-58 to G-234; E-59 to G-234; P-60 to G-234; E-61 to G-234; P-62 to G-234; E-63 to G-234; G-64 to G-234; G-65 to G-234; R-66 to G-234; S-67 to G-234; R-68 to G-234; A-69 to G-234; G-70 to G-234; A-71 to

G-234; Q-72 to G-234; N-73 to G-234; D-74 to G-234; Q-75 to G-234; L-76 to
G-234; S-77 to G-234; T-78 to G-234; G-79 to G-234; P-80 to G-234; R-81 to
G-234; A-82 to G-234; A-83 to G-234; P-84 to G-234; E-85 to G-234; E-86 to
G-234; A-87 to G-234; E-88 to G-234; T-89 to G-234; L-90 to G-234; A-91 to
5 G-234; E-92 to G-234; T-93 to G-234; E-94 to G-234; P-95 to G-234; E-96 to
G-234; R-97 to G-234; H-98 to G-234; L-99 to G-234; G-100 to G-234; S-101
to G-234; Y-102 to G-234; L-103 to G-234; L-104 to G-234; D-105 to G-234;
S-106 to G-234; E-107 to G-234; N-108 to G-234; T-109 to G-234; S-110 to
G-234; G-111 to G-234; A-112 to G-234; L-113 to G-234; P-114 to G-234;
10 R-115 to G-234; L-116 to G-234; P-117 to G-234; Q-118 to G-234; T-119 to
G-234; P-120 to G-234; K-121 to G-234; Q-122 to G-234; P-123 to G-234;
Q-124 to G-234; K-125 to G-234; R-126 to G-234; S-127 to G-234; R-128 to
G-234; A-129 to G-234; A-130 to G-234; F-131 to G-234; S-132 to G-234;
H-133 to G-234; T-134 to G-234; Q-135 to G-234; V-136 to G-234; I-137 to
15 G-234; E-138 to G-234; L-139 to G-234; E-140 to G-234; R-141 to G-234;
K-142 to G-234; F-143 to G-234; S-144 to G-234; H-145 to G-234; Q-146 to
G-234; K-147 to G-234; Y-148 to G-234; L-149 to G-234; S-150 to G-234;
A-151 to G-234; P-152 to G-234; E-153 to G-234; R-154 to G-234; A-155 to
G-234; H-156 to G-234; L-157 to G-234; A-158 to G-234; K-159 to G-234;
20 N-160 to G-234; L-161 to G-234; K-162 to G-234; L-163 to G-234; T-164 to
G-234; E-165 to G-234; T-166 to G-234; Q-167 to G-234; V-168 to G-234;
K-169 to G-234; I-170 to G-234; W-171 to G-234; F-172 to G-234; Q-173 to
G-234; N-174 to G-234; R-175 to G-234; R-176 to G-234; Y-177 to G-234;
K-178 to G-234; T-179 to G-234; K-180 to G-234; R-181 to G-234; K-182 to
25 G-234; Q-183 to G-234; L-184 to G-234; S-185 to G-234; S-186 to G-234;
E-187 to G-234; L-188 to G-234; G-189 to G-234; D-190 to G-234; L-191 to
G-234; E-192 to G-234; K-193 to G-234; H-194 to G-234; S-195 to G-234;
S-196 to G-234; L-197 to G-234; P-198 to G-234; A-199 to G-234; L-200 to
G-234; K-201 to G-234; E-202 to G-234; E-203 to G-234; A-204 to G-234;
30 F-205 to G-234; S-206 to G-234; R-207 to G-234; A-208 to G-234; S-209 to

G-234; L-210 to G-234; V-211 to G-234; S-212 to G-234; V-213 to G-234; Y-214 to G-234; N-215 to G-234; S-216 to G-234; Y-217 to G-234; P-218 to G-234; Y-219 to G-234; Y-220 to G-234; P-221 to G-234; Y-222 to G-234; L-223 to G-234; Y-224 to G-234; C-225 to G-234; V-226 to G-234; G-227 to G-234; S-228 to G-234; W-229 to G-234; of SEQ ID NO:2. Polynucleotides encoding these polypeptides are also encompassed by the invention.

Further embodiments of the invention are directed to C-terminal deletions of the NKX3 polypeptides of the invention, described by the general formula 1 to n, where n is any one of the integers from 2 to 233 corresponding to the position of amino acid residue identified in SEQ ID NO:2, and preferably corresponds to a residue identified in one of the C-terminal deletions specified herein. In specific embodiments, C-terminal deletions of the NKX3 polypeptides of the invention comprise, or alternatively, consist of, amino acid residues: M-1 to F-233; M-1 to A-232; M-1 to P-231; M-1 to S-230; M-1 to W-229; M-1 to S-228; M-1 to G-227; M-1 to V-226; M-1 to C-225; M-1 to Y-224; M-1 to L-223; M-1 to Y-222; M-1 to P-221; M-1 to Y-220; M-1 to Y-219; M-1 to P-218; M-1 to Y-217; M-1 to S-216; M-1 to N-215; M-1 to Y-214; M-1 to V-213; M-1 to S-212; M-1 to V-211; M-1 to L-210; M-1 to S-209; M-1 to A-208; M-1 to R-207; M-1 to S-206; M-1 to F-205; M-1 to A-204; M-1 to E-203; M-1 to E-202; M-1 to K-201; M-1 to L-200; M-1 to A-199; M-1 to P-198; M-1 to L-197; M-1 to S-196; M-1 to S-195; M-1 to H-194; M-1 to K-193; M-1 to E-192; M-1 to L-191; M-1 to D-190; M-1 to G-189; M-1 to L-188; M-1 to E-187; M-1 to S-186; M-1 to S-185; M-1 to L-184; M-1 to Q-183; M-1 to K-182; M-1 to R-181; M-1 to K-180; M-1 to T-179; M-1 to K-178; M-1 to Y-177; M-1 to R-176; M-1 to R-175; M-1 to N-174; M-1 to Q-173; M-1 to F-172; M-1 to W-171; M-1 to I-170; M-1 to K-169; M-1 to V-168; M-1 to Q-167; M-1 to T-166; M-1 to E-165; M-1 to T-164; M-1 to L-163; M-1 to K-162; M-1 to L-161; M-1 to N-160; M-1 to K-159; M-1 to A-158; M-1 to L-157; M-1 to H-156; M-1 to A-155; M-1 to R-154; M-1 to E-153; M-1 to P-152; M-1 to A-151; M-1 to S-150; M-1 to L-149; M-1 to Y-148; M-1 to

K-147; M-1 to Q-146; M-1 to H-145; M-1 to S-144; M-1 to F-143; M-1 to K-142; M-1 to R-141; M-1 to E-140; M-1 to L-139; M-1 to E-138; M-1 to I-137; M-1 to V-136; M-1 to Q-135; M-1 to T-134; M-1 to H-133; M-1 to S-132; M-1 to F-131; M-1 to A-130; M-1 to A-129; M-1 to R-128; M-1 to S-127; M-1 to R-126; M-1 to K-125; M-1 to Q-124; M-1 to P-123; M-1 to Q-122; M-1 to K-121; M-1 to P-120; M-1 to T-119; M-1 to Q-118; M-1 to P-117; M-1 to L-116; M-1 to R-115; M-1 to P-114; M-1 to L-113; M-1 to A-112; M-1 to G-111; M-1 to S-110; M-1 to T-109; M-1 to N-108; M-1 to E-107; M-1 to S-106; M-1 to D-105; M-1 to L-104; M-1 to L-103; M-1 to Y-102; M-1 to S-101; M-1 to G-100; M-1 to L-99; M-1 to H-98; M-1 to R-97; M-1 to E-96; M-1 to P-95; M-1 to E-94; M-1 to T-93; M-1 to E-92; M-1 to A-91; M-1 to L-90; M-1 to T-89; M-1 to E-88; M-1 to A-87; M-1 to E-86; M-1 to E-85; M-1 to P-84; M-1 to A-83; M-1 to A-82; M-1 to R-81; M-1 to P-80; M-1 to G-79; M-1 to T-78; M-1 to S-77; M-1 to L-76; M-1 to Q-75; M-1 to D-74; M-1 to N-73; M-1 to Q-72; M-1 to A-71; M-1 to G-70; M-1 to A-69; M-1 to R-68; M-1 to S-67; M-1 to R-66; M-1 to G-65; M-1 to G-64; M-1 to E-63; M-1 to P-62; M-1 to E-61; M-1 to P-60; M-1 to E-59; M-1 to P-58; M-1 to E-57; M-1 to P-56; M-1 to E-55; M-1 to P-54; M-1 to D-53; M-1 to R-52; M-1 to Q-51; M-1 to R-50; M-1 to Q-49; M-1 to S-48; M-1 to S-47; M-1 to T-46; M-1 to R-45; M-1 to G-44; M-1 to G-43; M-1 to Q-42; M-1 to R-41; M-1 to Q-40; M-1 to A-39; M-1 to G-38; M-1 to D-37; M-1 to R-36; M-1 to L-35; M-1 to I-34; M-1 to D-33; M-1 to Q-32; M-1 to I-31; M-1 to L-30; M-1 to F-29; M-1 to S-28; M-1 to T-27; M-1 to L-26; M-1 to P-25; M-1 to K-24; M-1 to S-23; M-1 to P-22; M-1 to T-21; M-1 to P-20; M-1 to P-19; M-1 to A-18; M-1 to A-17; M-1 to G-16; M-1 to E-15; M-1 to A-14; M-1 to K-13; M-1 to A-12; M-1 to E-11; M-1 to G-10; M-1 to P-9; M-1 to R-8; M-1 to P-7; M-1 to E-6; of SEQ ID NO:2. Polynucleotides encoding these polypeptides are also encompassed by the invention.

Further embodiments of the invention are directed to polypeptide fragments comprising, or alternatively, consisting of, amino acid residues

described by the general formula m to n, where m and n correspond to any one of the amino acid residues specified above for these symbols, respectively. Polynucleotides encoding these polypeptides are also encompassed by the invention.

5 ***NKX3.1 is a Homeoprotein and has a Role in Prostate Function***

10 NKX3.1 and its mouse counterpart represent the first mammalian homologues of the drosophila NK-3 gene. The human and mouse proteins are highly homologous, sharing 100 amino acid identity within the homeodomain regions, and are also identical to the drosophila gene within the homeodomain helix III and nearly identical within the N-terminal arm, regions which directly contact DNA and define DNA binding specificity in other homeodomains (Gehring
15 *et al.*, *Annu. Rev. Biochem.* 63:487-526 (1994)). Conservation of a tyrosine at position 54, which is conserved in the mouse and human homeodomains, largely defines the unique DNA binding specificity for a "CAAG" binding domain seen with NK family members, as opposed to the "TAAT" core which is preferentially
20 bound by most other the homeodomain proteins (See, reviews in Harvey, *Dev. Biol.* 178:203-216 (1996)). NKX3.1 has NK family-like DNA binding properties in that the protein has binds preferentially to a "CAAG" DNA core, confirming that NKX3.1 similar to other NK family proteins in its DNA binding properties.
25 (See, Example 7). The fact that the presumed DNA binding regions of the human and mouse homeodomain are essentially identical with the same region of drosophila NK-3, but not with NK-2 or NK4, may reflect a yet more refined DNA binding specificity that is unique to the NK-3-like homeodomains. Since it appears that NKX3.1 plays a central role in transcriptional regulation of prostate function (See, Examples below), this would provide an obvious mechanism for targeting transcriptional regulation to a unique set of genes as compared to those regulated by other NK-like proteins.

Outside the homeodomain the mouse and human gene sequences are also highly conserved but do not have strong homology with drosophila NK-3. Most intriguing is a C-terminal stretch of amino acids that is highly conserved between the mouse and human proteins and is exceptionally rich in tyrosine and serine residues. The fact that these are potential sites for post-transnational phosphorolation and that they are strongly conserved between that mouse and human suggests that they this region may play a role in regulating the function of the protein.

The highly restricted expression of NKX3.1 and the observed androgen regulation (*See*, Example 5 below), suggests that the gene may play a role in androgen-driven differentiation of prostatic and/or other urogenital tissues. Studies on the mouse NKX3.1 gene indicate that his gene is a strong candidate for playing a central role in maintenance of the prostatic phenotype in adults as well as its establishment during prostate development. NKX3.1 expression in mouse embryos during late gestation coincides with the beginning of prostatic bud formation, and *situ* localization studies indicate expression at this stage is restricted to epithelial cells within the prostatic buds, urogenital sinus, and testis. (*See*, Example 9 below) NKX3.1 is also expressed at and different stages and at lower levels in epithelial cell types of several other tissues (*See*, Examples 9 and 10 below) indicating a potentially broader role in epithelial cell development.

Embryonic expression of NKX3.1 occurs at a time when the prostatic epithelium is not androgen-responsive, indicating that expresison of the gene during early development is not androgen dependent. However, later in development there is a surge in NKX3.1 expression which parallels the androgen-driven maturation of adult prostate and seminal vesicle expression (*See*, Example 9 below). These data are consistent with the fact that human NKX3.1 expression was only found in prostate, testis, and an androgen-dependent prostatic carcinoma line. In addition, a dramatic and immediate loss of expression was seen following upon castration of adult mice, which is again consistent with the requirement of androgens for NKX3.1 in LNCaP cells (and the absence of expression in prostatic

cell lines which have lost androgen responsiveness). Taken together these studies suggest that NKX3.1 and its mouse counterpart, in addition to participating in early urogenital development, may play a role in the androgen-driven maintenance of the differentiated state of prostatic tissue.

5 ***Cancer Diagnosis and Prognosis***

An intriguing observation is the fact that this new prostate-specific gene, NKX3.1, maps to a chromosomal locus whose loss is implicated in the progression of prostate cancer (*See*, Example 6, below) and may well be involved in the progression of other forms of cancer. In prostate cancer there is a transition from androgen-dependent growth to androgen-independent growth which is correlated with a loss of prostate tissue phenotype. Although, many studies have addressed this switch to androgen-independence, including several using both the androgen-dependent cell line, LNCaP, and androgen-independent cell lines, PC-3 and DU-145, the underlying mechanisms causing this fundamental transition remain unknown.

Studies have focused on cytogenetic changes and loss of heterozygosity (LOH) studies to identify genetic changes occurring during the progression of prostate cancer and have identified a region spanning 8p21-22 which is thought to contain at least two genes that are lost during the progression of prostate cancer. Detailed deletion mapping has defined a 1.2 Mb commonly deleted within bands 8p22-p21.3, while a second region of common deletion was identified between at 8p21-8p11.22, suggesting that at least 2 tumor suppresser genes are present on chromosome 8p (Suzuki *et al.*, *Genes, Chromosomes, and Cancer* 13:168-174 (1995)). The distal deleted region overlaps with the region commonly deleted in hepatocellular carcinomas, colorectal cancers and non-small cell lung cancers (Emi *et al.*, *Genomics* 15:530-534 (1992); Fujiwara *et al.*, *Genes, Chromosomes, and Cancer* 10:7-14 (1994)). The proximal region, in which the NKX3.1 gene maps, is deleted in up 60-80% prostate cancers (Suzuki, *et al.*,

Genes, Chromosomes and Cancer 13:168-174 (1995), Bova *et al.*, *Cancer Res.* 53:3869-3873 (1993), MacGrogan *et al.*, *Genes, Chromosomes and Cancer* 10:151-159 (1994), Trapman *et al.*, *Cancer Res.* 54:6061-6064 (1994), Macoska, *et al.*, *Cancer Res.* 55:5390-5395 (1995), and Vocke *et al.*, *Cancer Res.* 56:2411-2416 (1996) and overlaps with a region that is commonly deleted in colorectal cancers (Fujiwara *et al.*, 1993). Recently, Vocke *et al.* (1996) using the most precise micro-dissection resources available for DNA analysis, found that the loss of markers within 8p12-21, is a very frequent event (62-75%) and is equally likely to occur in early and advanced disease. They interpret these data to indicate that loss at 8p12-21 may be a seminal early event in tumor development, likely prior to loss of androgen-responsiveness and associated loss of the differentiated prostatic phenotype.

Since the NKX3.1 gene appears to be located on 8p21 (*see*, Example 6 below), the data are consistent with a role for the gene in androgen-driven maintenance of prostate tissue phenotype.

Thus, it is believed that certain tissues in mammals with prostate cancer express significantly decreased levels of the NKX3.1 protein and mRNA encoding the NKX3.1 protein when compared to a corresponding "standard" mammal, i.e., a mammal of the same species not having the cancer. Further, it is believed that decreased levels of the NKX3.1 protein can be detected in certain body fluids (e.g., sera, plasma, urine, and spinal fluid) from mammals with prostate cancer when compared to sera from mammals of the same species not having the cancer. Thus, the invention provides a diagnostic method useful during prostate as well as other cancer diagnosis, which involves assaying the expression level of the gene encoding the NKX3.1 protein in mammalian cells or body fluid and comparing the gene expression level with a standard NKX3.1 gene expression level, whereby a decrease in the gene expression level over the standard is indicative of prostate tumors and other cancers.

Furthermore, the NKX3.1 gene could be used in PCR or other DNA based tests well known to those of skill in the art for mutations and/or loss of this gene

during the progression of cancer which could be used diagnostically to monitor the progression and potential medication responsiveness to prostate and other forms of cancer. Similarly, an antibody directed against the NKX3.1 protein or mutant forms of the protein could be used as a marker to monitor the progression and potential medication responsiveness to prostate cancer using immunoassays well known to those of skill the art.

Where a tumor diagnosis has already been made according to conventional methods, the present invention is useful as a prognostic indicator, whereby patients exhibiting decreased NKX3.1 gene expression will experience a worse clinical outcome relative to patients expressing the gene at an enhanced level.

By "assaying the expression level of the gene encoding the NKX3.1 protein" is intended qualitatively or quantitatively measuring or estimating the level of the NKX3.1 protein or the level of the mRNA encoding the NKX3.1 protein in a first biological sample either directly (e.g., by determining or estimating absolute protein level or mRNA level) or relatively (e.g., by comparing to the NKX3.1 protein level or mRNA level in a second biological sample).

Preferably, the NKX3.1 protein level or mRNA level in the first biological sample is measured or estimated and compared to a standard NKX3.1 protein level or mRNA level, the standard being taken from a second biological sample obtained from an individual not having the cancer. As will be appreciated in the art, once a standard NKX3.1 protein level or mRNA level is known, it can be used repeatedly as a standard for comparison.

By "biological sample" is intended any biological sample obtained from an individual, cell line, tissue culture, or other source which contains NKX3.1 protein or mRNA. Biological samples include mammalian body fluids (such as sera, plasma, urine, synovial fluid and spinal fluid) which contain NKX3.1 protein, and ovarian, prostate, heart, placenta, pancreas liver, spleen, lung, breast and umbilical tissue.

The present invention is useful for detecting cancer in mammals. In particular the invention is useful during diagnosis of prostate cancers in mammals.

Preferred mammals include monkeys, apes, cats, dogs, cows, pigs, horses, rabbits and humans. Particularly preferred are humans.

Total cellular RNA can be isolated from a biological sample using the single-step guanidinium-thiocyanate-phenol-chloroform method described in Chomczynski and Sacchi, *Anal. Biochem.* 162:156-159 (1987). Levels of mRNA encoding the NKX3.1 protein are then assayed using any appropriate method. These include Northern blot analysis (Harada *et al.*, *Cell* 63:303-312 (1990)), S1 nuclease mapping (Fujita *et al.*, *Cell* 49:357- 367 (1987)), the polymerase chain reaction (PCR), reverse transcription in combination with the polymerase chain reaction (RT-PCR) (Makino *et al.*, *Technique* 2:295-301 (1990)), and reverse transcription in combination with the ligase chain reaction (RT-LCR).

Assaying NKX3.1 protein levels in a biological sample can occur using antibody-based techniques. For example, NKX3.1 protein expression in tissues can be studied with classical immunohistological methods (Jalkanen, M., *et al.*, *J. Cell. Biol.* 101:976-985 (1985); Jalkanen, M., *et al.*, *J. Cell . Biol.* 105:3087-3096 (1987)).

Other antibody-based methods useful for detecting NKX3.1 protein gene expression include immunoassays, such as the enzyme linked immunosorbent assay (ELISA) and the radioimmunoassay (RIA).

Suitable labels are known in the art and include enzyme labels, such as, Glucose oxidase, and radioisotopes, such as iodine (^{125}I , ^{121}I), carbon (^{14}C), sulfur (^{35}S), tritium (^3H), indium (^{112}In), and technetium ($^{99\text{m}}\text{Tc}$), and fluorescent labels, such as fluorescein and rhodamine, and biotin.

Therapeutics

Since it appears that the loss of NKX3.1 activity plays a role in the loss of differentiated prostatic phenotype in prostate cancer, administration of this protein, or an NKX3.1 agonist, could be used to prevent or treat the loss of differentiated prostate tissue, and thereby prevent or treat prostate cancer. In

addition, administration of this protein, or an NKX3.1 agonist, could be used to prevent or treat other forms of cancer.

Modes of administration

5 It will be appreciated that conditions caused by a decrease in the standard or normal level of NKX3.1 activity in an individual, can be treated by administration of NKX3.1 protein, or an NKX3.1 agonist. Thus, the invention further provides a method of treating an individual in need of an increased level of NKX3.1 activity comprising administering to such an individual a pharmaceutical composition comprising an effective amount of an isolated
10 NKX3.1 polypeptide of the invention, or an NKX3.1 agonist, effective to increase the NKX3.1 activity level in such an individual.

As a general proposition, the total pharmaceutically effective amount of NKX3.1 polypeptide administered parenterally per dose will be in the range of about 1 µg/kg/day to 10 mg/kg/day of patient body weight, although, as noted
15 above, this will be subject to therapeutic discretion. More preferably, this dose is at least 0.01 mg/kg/day, and most preferably for humans between about 0.01 and 1 mg/kg/day for the hormone. If given continuously, the NKX3.1 polypeptide is typically administered at a dose rate of about 1 µg/kg/hour to about 50 µg/kg/hour, either by 1-4 injections per day or by continuous subcutaneous
20 infusions, for example, using a mini-pump. An intravenous bag solution may also be employed.

Pharmaceutical compositions containing the NKX3.1 of the invention may be administered orally, rectally, parenterally, intracisternally, intravaginally, intraperitoneally, topically (as by powders, ointments, drops or transdermal patch),
25 buccally, or as an oral or nasal spray. By "pharmaceutically acceptable carrier" is meant a non-toxic solid, semisolid or liquid filler, diluent, encapsulating material or formulation auxiliary of any type. The term "parenteral" as used herein refers

to modes of administration which include intravenous, intramuscular, intraperitoneal, intrasternal, subcutaneous and intraarticular injection and infusion.

Selection of Compounds Capable of Regulating Expression of NKX3.1

As the invention also includes isolated genomic DNA molecules comprising the 5' flanking region of NKX3.1, including the promoter for these splice variants, yet another aspect of the invention is related to a method for identifying compounds capable of enhancing or inhibiting expression of NKX3.1. In order to determine the effect of such compounds, reporter plasmids are constructed by linking a portion of the DNA located 5' to the transcription start site of any of NKX3.1 in front of a reporter gene. Such constructs are then transfected into appropriate cell lines. Compounds that are to be tested for their ability to increase or decrease expression from the NKX3.1 promoter are then administered to the cell bearing the reporter construct, and the effect of each compound on reporter gene expression is determined by comparing that level of expression to the expression level in a control cell bearing the reporter construct, where the test compound has not been administered to the control cell.

The DNA sequence of the 5' flanking region of the NKX3.1 gene is shown in Figure 5 (SEQ ID NO:8) (ATCC No. 209006). Of course, since the nucleotide sequence is known, routine methods are available for producing such nucleic acid molecules synthetically (see, for example, *Synthesis and Application of DNA and RNA*, S.A. Narang, ed., 1987, Academic Press, San Diego, CA). Alternatively, such isolated nucleic acid molecules of the present invention can be generated as follows. The NKX3.1 gene promoter region is obtained by amplification using the polymerase chain reaction (PCR). The amplified fragment is then inserted into an appropriate plasmid (such as, for example, pCAT™ (Promega, Madison, WI)). Nested deletion plasmids are then generated using the commercially available "Erase-a-Base" System (Promega, Madison, WI) as described in Henikoff, *Gene* 28:351-359 (1984)). Thus, only routine experimentation would be required to

generate any of the isolated nucleic acid molecules of the present invention which are capable of enhancing or inhibiting gene expression.

The nucleic acid molecules of the present invention can include the NKX3.1 promoter and *cis*-acting enhancer and/or silencer elements capable of affecting gene transcription. For simplicity, these isolated nucleic acid molecules of the present invention are referred to below as "NKX3.1 transcriptional regulatory elements" or "transcriptional elements." As indicated, to determine the effect of a transcriptional element of the present invention on gene expression, nested deletion reporter plasmids can be generated containing a transcriptional element of the present invention linked in front of the chloramphenicol acetyltransferase (CAT) reporter gene. Such recombinant DNA molecules of the present invention actually generated by the inventors include transcriptional elements inserted, in both orientations, into the *Xba*I site of pBLCAT2 vector (Luckow, B., Schütz, G., *Nucleic Acids Res.* 15:5490 (1987)).

By the invention, a recombinant DNA molecule containing a transcriptional element of the present invention is used to transiently transfect an appropriate cell line such as, for example, human choriocarcinoma cell lines (JEG-3 and JAR), the human prostate carcinoma cell line PC-3, or the monkey kidney cell line CV-1, all of which are available from the American Type Culture Collection. In addition to using the CAT system for reporter gene analyses, the hGH transient expression system can also be used (Selden *et al.*, *Mol. Cell Biol.* 6:3173-3179 (1986)) or other systems that are based on the expression of β -galactosidase (An *et al.*, *Mol. Cell Biol.* 2:1628-1632 (1982)) and xanthine-guanine phosphoribosyl transferase (Chu *et al.*, *Nucleic Acids Res.* 13:2921-2930 (1985)).

A transcriptional element of the present invention may be inserted into an appropriate vector in accordance with conventional techniques, including blunt-ending or staggered-ending termini for ligation, restriction enzyme digestion to provide appropriate termini, filling in of cohesive ends as appropriate, alkaline phosphatase treatment to avoid undesirable joining, and ligation with appropriate

ligases. Techniques for such manipulations are disclosed by Maniatis, T., *et al.*, *infra*, and are well known in the art. Clones containing a transcriptional element of the present invention may be identified by any means which specifically selects for a NKX3.1 enhancer or silencer region DNA such as, for example by hybridization with an appropriate nucleic acid probe(s) containing a sequence complementary to all or part of the transcriptional element. Oligonucleotide probes specific for a transcriptional element of the present invention can be designed simply by reference to the sequences disclosed in Figure 5. Techniques for nucleic acid hybridization and clone identification are disclosed by Maniatis, T., *et al.*, (In: *Molecular Cloning, A Laboratory Manual*, Cold Spring Harbor Laboratories, Cold Spring Harbor, NY (1982)), and by Hames, B.D., *et al.*, (In: *Nucleic Acid Hybridization, A Practical Approach*, IRL Press, Washington, DC (1985)). To facilitate the detection of the desired clone containing a transcriptional element of the present invention, the above-described nucleic acid probe may be labeled with a detectable group. Such detectable groups can be any material having a detectable physical or chemical property. Such materials have been well-developed in the field of nucleic acid hybridization and in general most any label useful in such methods can be applied to the present invention. Particularly useful are radioactive labels, such as ^{32}P , ^3H , ^{14}C , ^{35}S , ^{125}I , or the like. Any radioactive label may be employed which provides for an adequate signal and has a sufficient half-life. The oligonucleotide may be radioactively labeled, for example, by "nick-translation" by well-known means, as described in, for example, Rigby, P.J.W., *et al.*, *J. Mol. Biol.* 113:237 (1977) and by T4 DNA polymerase replacement synthesis as described in, for example, Deen, K.C., *et al.*, *Anal. Biochem.* 135:456 (1983). Alternatively, polynucleotides are also useful as nucleic acid hybridization probes when labeled with a non-radioactive marker such as biotin, an enzyme or a fluorescent group. See, for example, Leary, J.J., *et al.*, *Proc. Natl. Acad. Sci. USA* 80:4045 (1983); Renz, M., *et al.*, *Nucl. Acids Res.* 12:3435 (1984); and Renz, M., *EMBO J.* 6:817 (1983).

As used herein, "heterologous protein" is intended to refer to a peptide sequence that is heterologous to the transcriptional regulatory elements of the invention. A skilled artisan will recognize that, if desired, the teaching herein will also apply to the expression of genetic sequences encoding the NKX3.1 protein, or splice variants thereof, by such transcriptional regulatory elements. The reporter genes for use in the screening assay described below can code for either the NKX3.1 protein, or splice variants thereof, or a heterologous protein. Alternatively, detection of reporter gene expression can be at the mRNA level, such as, for example, detection of NKX3.1 mRNA.

To express a reporter gene under the control of the transcriptional regulatory elements of the invention, the gene must be "operably-linked" to the regulatory element. An operable linkage is a linkage in which a desired sequence is connected to a transcriptional or translational regulatory sequence (or sequences) in such a way as to place expression (or operation) of the desired sequence under the influence or control of the regulatory sequence.

Two DNA sequences (such as a reporter gene and a promoter region sequence linked to the 5' end of the reporter gene) are said to be operably linked if induction of promoter function results in the transcription of the reporter gene and if the nature of the linkage between the two DNA sequences does not (1) result in the introduction of a frame-shift mutation (if reporter protein activity is necessary for detection of reporter gene expression), (2) interfere with the ability of the expression regulatory sequences to direct reporter gene expression, or (3) interfere with the ability of reporter gene to be transcribed by the promoter region sequence. Thus, a promoter would be operably linked to a DNA sequence if the promoter were capable of affecting transcription of that DNA sequence.

In a similar manner, a transcriptional regulatory element of the present invention that enhances or represses gene expression may be operably-linked to such a promoter. Exact placement of the element in the nucleotide chain is not critical as long as the element is located at a position from which the desired effects on the operably linked promoter may be revealed. A nucleic acid molecule,

such as DNA, is said to be "capable of expressing" a polypeptide if it contains expression control sequences which contain transcriptional regulatory information and such sequences are operably linked to the nucleotide sequence which encodes the polypeptide. For the complete control of gene expression, all transcriptional and translational regulatory elements (or signals) that are operably linked to a heterologous gene should be recognizable by the appropriate host. By "recognizable" in a host is meant that such signals are functional in such host.

The NKX3.1 transcriptional regulatory elements of the present invention, obtained through the methods described above, and preferably in a double-stranded form, may be operably linked to a heterologous gene (such as a reporter gene), preferably in an expression vector, and introduced into a host cell, preferably a eukaryotic cell, to assay reporter gene expression. Preferred eukaryotic cells include prostate cell lines, choriocarcinoma cell lines, breast cancer cell lines, prostate carcinoma cell lines and kidney cell lines.

As is widely known, translation of eukaryotic mRNA is initiated at the codon that encodes the first methionine. For this reason, it is preferable to ensure that the linkage between a eukaryotic promoter and a reporter gene does not contain any intervening codons that are capable of encoding a methionine. The presence of such codons results either in a formation of a fusion protein (if the AUG codon is in the same reading frame as the DNA encoding the heterologous protein) or a frame-shift mutation (if the AUG codon is not in the same reading frame as the reporter gene).

If desired, a fusion product of a reporter protein may be constructed. For example, the sequence coding for the reporter protein may be linked to a signal sequence which will allow secretion of the protein from, or the compartmentalization of the protein in, a particular host. Such signal sequences may be designed with or without specific protease sites such that the signal peptide sequence is amenable to subsequent removal. Alternatively, the native signal sequence for this protein may be used.

The transcriptional regulatory elements of the invention can be selected to allow for repression or activation, so that expression of the operably linked reporter genes can be modulated. Translational signals are not necessary when it is desired to express antisense RNA sequences or to assay reporter gene expression via mRNA detection.

If desired, the non-transcribed and/or non-translated regions 3' to the reporter gene can be obtained by the above-described cloning methods. The 3'-non-transcribed region may be retained for its transcriptional termination regulatory sequence elements; the 3'-non-translated region may be retained for its translational termination regulatory sequence elements, or for those elements that direct polyadenylation in eukaryotic cells. Where the native expression control sequences signals do not function satisfactorily host cell, then sequences functional in the host cell may be substituted.

To transform a mammalian cell with the DNA constructs of the invention many vector systems are available, depending upon whether it is desired to insert the reporter gene product into the host cell chromosomal DNA, or to allow it to exist in an extrachromosomal form. If the reporter gene and an operably linked promoter are introduced into a recipient eukaryotic cell as a non-replicating DNA (or RNA) molecule, which may either be a linear molecule or, more preferably, a closed covalent circular molecule that is incapable of autonomous replication, reporter gene expression may occur through the transient expression of the introduced sequence.

Genetically stable transformants may be constructed with vector systems, or transformation systems, whereby the reporter gene is integrated into the host chromosome. Such integration may occur *de novo* within the cell or, in a most preferred embodiment, be assisted by transformation with a vector that functionally inserts itself into the host chromosome. Vectors capable of chromosomal insertion include, for example, retroviral vectors, transposons or other DNA elements which promote integration of DNA sequences in

chromosomes, especially DNA sequence homologous to a desired chromosomal insertion site.

Cells that have stably integrated the introduced DNA into their chromosomes are selected by also introducing one or more markers that allow for selection of host cells which contain the desired sequence. For example, the marker may provide biocide resistance, e.g., resistance to antibiotics, or heavy metals, such as copper, or the like. The selectable marker gene can either be directly linked to the reporter gene, or introduced into the same cell by co-transfection. In another embodiment, the introduced sequence is incorporated into a plasmid or viral vector capable of autonomous replication in the recipient host. Any of a wide variety of vectors may be employed for this purpose, as outlined below. Factors of importance in selecting a particular plasmid or viral vector include: the ease with which recipient cells that contain the vector may be recognized and selected from those recipient cells which do not contain the vector; the number of copies of the vector which are desired in a particular host; and whether it is desirable to be able to "shuttle" the vector between host cells of different species.

Preferred eukaryotic plasmids include those derived from the bovine papilloma virus, vaccinia virus, and SV40. Such plasmids are well known in the art and are commonly or commercially available. For example, mammalian expression vector systems in which it is possible to cotransfect with a helper virus to amplify plasmid copy number, and, integrate the plasmid into the chromosomes of host cells have been described (Perkins, A.S. *et al.*, *Mol. Cell Biol.* 3:1123 (1983); Clontech, Palo Alto, California). Particularly preferred are vectors derived from pCAT-Basic, pCAT-Enhancer and pCAT-Promoter vectors (Promega, Madison, WI).

Once the vector or DNA sequence containing the construct(s) is prepared for expression, the DNA construct(s) is introduced into an appropriate host cell by any of a variety of suitable means, including transfection, electroporation or delivery by liposomes. DEAE dextran, calcium phosphate, and preferably, the transfection reagent DOTAP, may be useful in the transfection protocol.

After the introduction of the vector *in vitro*, recipient cells are grown in a selective medium, that is, medium that selects for the growth of vector-containing cells. Expression of the reporter gene results in the production mRNA and, if desired, reporter protein. According to the invention, this expression can
5 take place in a continuous manner in the transformed cells, or in a controlled manner. If desired, in *in vitro* culture, the reporter protein is isolated and purified in accordance with conventional conditions, such as extraction, precipitation, chromatography, affinity chromatography, electrophoresis, or the like. Alternatively, levels of reporter protein expression can be assayed according to
10 conventional protein assays, such as, for example, the CAT expression system.

The NKX3.1 transcriptional regulatory elements of the present invention (i.e., the NKX3.1 promoter, as well as isolated nucleic acid molecules capable of enhancing and/or repressing gene expression) are useful for screening drugs, ligands and/or other *trans*-acting agents to determine which are capable of
15 affecting expression of NKX3.1 or any splice variant thereof. By the invention, *trans*-acting factors can be identified by their ability to up-regulate or down-regulate NKX3.1 expression. As used herein, by "NKX3.1 *trans*-acting agent" is intended a drug, ligand, or other compound capable interacting, either directly or indirectly, with a NKX3.1 transcriptional regulatory element of the present
20 invention to enhance or repress gene expression. Such NKX3.1 *trans*-acting elements which interact directly with a transcriptional regulatory element of the present invention include those, which, for example, bind directly to the element and either enhance or repress gene expression. NKX3.1 *trans*-acting agents which interact indirectly with a transcriptional regulatory element of the present invention
25 include those which, for example, bind to and induce activity of a second *trans*-acting agent (e.g., a receptor molecule) which itself then, either alone or complexed to the first *trans*-acting agent, binds to the element and either enhances or represses gene expression. One type of *trans*-acting agent is a triplex-forming oligonucleotide. Administration of a suitable oligonucleotide will result in the
30 formation of a triple helix between the oligonucleotide and the NKX3.1 promoter,

which will inhibit transcription from that promoter (Ebbinghaus, S.W. *et al.*, *Gene Therapy* 3: 287-297 (1996); Roy, C., *Eur. J. Biochem.* 220: 493-503 (1994)). Because the genomic sequence of the region 5' of the NKX3.1 gene is given herein (See FIG. 5 and SEQ ID NO:8), one of ordinary skill in the art will readily be able to design suitable oligonucleotides (also called "anti-sense" oligonucleotides) which can inhibit expression from the NKX3.1 promoter. One region which is especially useful for anti-sense design is the 5' untranslated region (*J. Biol. Chem.* 266: 18162-18171 (1991)), which of course is not included in a cDNA, but is included in the genomic sequence disclosed herein.

Thus, in one aspect, the invention provides a screening assay for determining whether any given compound is capable of up-regulating or down-regulating expression from the NKX3.1 promoter, leading to an increase or decrease of NKX3.1 production.

The screening assay involves (1) providing a host cell transfected with a recombinant nucleic acid molecule containing a NKX3.1 transcriptional regulatory element of the present invention and a reporter gene, wherein the transcriptional element is operably linked to the reporter gene; (2) administering a candidate NKX3.1 *trans*-acting agent to the transfected host cell; and (3) determining the effect on reporter gene expression.

In a preferred embodiment, the invention provides a screening assay for the identification of substances capable of altering the expression from the NKX3.1 promoter, comprising:

(a) measuring the level of expression of a reporter gene in a test cell, wherein said test cell is transformed with a recombinant DNA molecule comprising a reporter gene operably linked to a DNA molecule comprising the promoter of NKX3.1, and wherein a candidate NKX3.1 *trans*-acting agent is administered to said test cell;

(b) measuring the level of expression of said reporter gene in a control cell, wherein said control cell is transformed with the recombinant DNA molecule of step (a); and

(c) comparing the level of expression of said reporter gene in said test cell to the level of said reporter gene in said control cell.

Suitable and preferred host cells, transfection methods, expression vectors, promoters, and reporter genes, are described above and will be known in the art.

5 ***Gene Therapy***

Gene therapy has been proposed as a method for treating disease states and genetic disorders that lack effective therapies. Gene therapy techniques can also be applied as a method to control expression of a protein and to assess its ability to modulate cellular events.

10 The genomic clone for the NKX3.1 gene of the present invention has been isolated (Figure 5). This clone contains the promoter region for the NKX3.1 gene which appears to be a prostate tissue specific promoter. Thus, the NKX3.1 promoter region could be useful as a prostate-specific DNA element to drive expression of any gene of interest in gene therapy techniques. Promoter elements
15 are shown in Figure 5.

Thus, for example, cells from a patient may be engineered with a polynucleotide, such as a DNA or RNA, encoding a polypeptide *ex vivo*, and the engineered cells then can be provided to a patient to be treated with a polypeptide. For example, cells may be engineered *ex vivo* by the use of a retroviral plasmid
20 vector containing RNA encoding a desired polypeptide. Such methods are well-known in the art and their use in the present invention will be apparent from the teachings herein.

Similarly, cells may be engineered *in vivo* for expression of a polypeptide *in vivo* by procedures well-known in the art. For example, a polynucleotide may
25 be engineered for expression in a replication defective retroviral vector, as discussed above. The retroviral expression construct then may be isolated and introduced into a packaging cell which is transduced with a retroviral plasmid vector containing RNA encoding a desired polypeptide such that packaging cell

now produces infectious viral particles contain the gene of interest. These producer cells may be administered to a patient for engineering cells *in vivo* and expression of the polypeptide *in vivo*. These and other methods for administering a polypeptide of interest by such methods will be apparent to those skilled in the art from the teachings of the present invention.

Retroviruses from which the retroviral plasmid vectors mentioned above may be derived include, but are not limited to, Moloney Murine Leukemia Virus, spleen necrosis virus, retroviruses such as Rous Sarcoma Virus, Harvey Sarcoma Virus, avian leukosis virus, gibbon ape leukemia virus, human immunodeficiency virus, adenovirus, Myeloproliferative Sarcoma Virus, and mouse mammary tumor virus. In one embodiment, the retroviral plasmid vector is derived from Moloney Murine Leukemia Virus.

The retroviral plasmid vector is employed to transduce packaging cell lines to form producer cell lines. Example of packaging cells which may be transfected include, but are not limited to, PE501, PA317, Y-2, Y-AM, PA12, T19-14X, VT-19-17-H2, YCRE, YCRIP, GP+E-86, GP+envAm12, and DAN cell lines as described in Miller A., *Human Gene Therapy 1*: 5-14 (1990). The vector may be transduced into the packaging cells through any means known in the art. Such means include, but are not limited to, electroporation, the use of liposomes, and CaPO_4 precipitation. In one alternative, the retroviral vector plasmid vector may be encapsulated into a liposome, or coupled to a lipid and then administered to a host.

The producer cell line will generate infectious retroviral vector particles, which include the nucleic acid sequence(s) encoding a desired polypeptide. Such retroviral particles then may be employed to transduce eukaryotic cells, either *in vitro* or *in vivo*. The transduced eukaryotic cells will express the nucleic acid sequence(s) encoding the polypeptide. Eukaryotic cells which may be transduced include, but are not limited to, embryonic stem cells, embryonic carcinoma cells, as well fibroblasts, epithelial cells and endothelial cells.

In addition to retroviral vectors, a variety of vectors have also been developed for gene delivery. These vectors derive from herpes simplex virus type 1 (HSV-1), adenovirus, adeno-associated virus (AAV) and retrovirus constructs (for review see Friedmann, T., *Trends Genet* 10:210-214 (1994); Jolly, D., *Cancer Gene Therapy* 1 (1994); Mulligan, R.C., *Science* 260:926-932 (1993); Smith, F. *et al.*, *Rest. Neurol. Neurosci.* 8:21-34 (1995)). Vectors based on HSV-1, including both recombinant virus vectors and amplicon vectors, as well as adenovirus vectors can assume an extrachromosomal state in the cell nucleus and mediate limited, long term gene expression. HSV-1 amplicon vectors can be grown to relatively high titers (10^7 transducing units/ml) and have the capacity to accommodate large fragments of foreign DNA (at least 15 kb, with 10 concatemeric copies per virion). AAV vectors (rAAV), available in comparable titers to amplicon vectors, can deliver genes (< 4.5 kb) to postmitotic, as well as mitotic cells in combination with adenovirus or herpes virus as helper virus. Long term transgene expression is achieved by replication and formation of "episomal" elements and/or through integration into the host cell genome at random or specific sites (for review see Samulski, R.J., *Current Opinion in Genetics and Development* 3:74-80 (1993); Muzyczka, N., *Curr. Top. Microbiol. Immunol.* 158:97-129 (1992)). HSV, adenovirus and rAAV vectors are all packaged in stable particles. Retrovirus vectors can accommodate 7-8 kb of foreign DNA and integrate into the host cell genome. Recent studies have demonstrated that elements from different viruses can be combined to increase the delivery capacity of vectors. For example, incorporation of elements of the HIV virion, including the matrix protein and integrase, into retrovirus vectors allows transgene cassettes to enter the nucleus of non-mitotic, as well as mitotic cells and potentially to integrate into the genome of these cells (Naldini, L. *et al.*, *Science* 272:263-267 (1996)); and inclusion of the vesicular stomatitis virus envelope glycoprotein (VSV-G) increases stability of retrovirus particles (Emi, N. *et al.*, *J. Virol.* 65:1202-1207 (1991)). As another example, inclusion of elements from Epstein Barr virus (EBV) - the DNA origin of replication, oriP, and the EBNA-1, within

HSV vectors allows nuclear replication of vectors in dividing human cells (Wang and Vos, in press).

Both HSV and AAV can deliver genes to dividing and non-dividing cells. In general, HSV virions are considered more highly infectious than AAV virions, with a ratio of virus particles: infectious units in the range of 10 for HSV (Browne, H. *et al.*, *J. Virol.* 70:4311-4316 (1996)) and up to thousands for AAV (Snyder, R.O. *et al.*, In Current Protocols in Human Genetics, Eds. Dracopoli, N. *et al.*, John Wiley and Sons: New York (1996), pp. 1-24), and both having a broad species range. These vectors deliver linear, double stranded DNA to the nucleus. If replication-competent HSV helper virus accompanies the vector DNA it commences replication as a rolling circle. If the helper virus is replication-defective or absent, the amplicon DNA is thought to exist as a linear extrachromosomal element without any defined episomal structure.

HSV-1 based vectors allow replication and packaging of DNA as a concatenate in HSV-1 virions in the presence of HSV-1 helper virus (Kwong, A.D. and Frenkel, N., *In Viral Vectors*, Eds. M.G. Kaplitt and A.D. Loewy, Academic Press: New York (1995), pp. 25-42). Vectors based on either HSV-1 or AAV can be generated using either HSV-1 or AAV helper virus packaging systems well known in the art. In addition, vectors can be generated free of helper virus using the helper virus-free packaging system described by Fraefel, C. *et al.*, *J. Virol.* 70: 7190-7197 (1996).

These gene therapy vectors, when encapsulated in either a herpesviral particle coat or an adeno-associated viral particle coat, capable of binding to a target host cell and introducing into the target host cell the polynucleotide sequence carried by the vector.

For the treatment of cancers and tumors, gene therapy can be used to introduce a gene into the tumor cells that expresses a protein which is toxic or can trigger a toxic effect against tumor cells. Genes for transfer into the neoplastic cells by the vectors are selected from those which target host cell usually by expression of a gene product in the host neoplastic cells. "Gene product" broadly

refers to proteins encoded by the particular gene. For the purposes of the invention, gene product also includes transcription products of the gene, particularly for use as antisense RNA. Genes are selected whose gene products serve to identify host cells, slow down or temporarily stimulate host cell growth in order to render the host cell more sensitive to chemotherapeutic agents and/or whose products target the host cell for cell death. Cell death can be accomplished by contacting the host cells, containing the gene product, with a subsequent treatment, either physical or chemical treatment. Alternatively, the gene products themselves may serve to kill the host cells or slow down cell growth. Such genes and gene products are known to those skilled in the art. The host cells targeted by the present hybrid vectors are those cells into which the hybrid vector infects and expresses the desired gene product and thus can constitute neoplastic cells infected by the hybrid vectors.

Useful gene products comprise: tumor suppressor genes, which encode transcription factors which suppress cell growth, such as the Rb gene for retinoblastoma or the p53 gene in colon cancer (Huang *et al.*, *Science* 242: 1563-1566 (1988); Barker, *et al.*, *Science* 249: 912-915 (1980); toxic proteins that are released by cells, such as a fusion protein comprising a toxin coupled to EGF ligand (Heinbrook *et al.*, *Proc. Natl. Acad. Sci. USA* 87: 4697 (1990)); products which themselves are capable of selective cell killing, such as anti-sense nucleic acid for essential cell proteins, such as replication proteins which serve to render the host cells incapable of further cell growth and division (Rosengberg *et al.*, *Nature* 313: 703-706 (1985); Preiss *et al.*, *Nature* 313:27-32 (1985) McGarry *et al.*, *Proc. Natl. Acad. Sci USA* 83: 399-403(1986); and prodrug activating genes such as thymidine kinase (Kramm *et al.*, *Brain Pathology* 5:345-381 (1995)).

Thus, one embodiment of the invention provides for a nucleic acid sequence encoding the polypeptide of interest contained in either the retroviral or amplicon vectors described above and placed under the control of the NKX3.1 promoter region of the present invention. Thus, such a vector having the NKX3.1 prostate-specific promoter element(s) operably linked to a gene of interest,

provides a vector that can specifically target the expression of such a gene to prostate tissue.

Chromosome Assays

5 The nucleic acid molecules of the present invention are also valuable for chromosome identification. The sequence is specifically targeted to and can hybridize with a particular location on an individual human chromosome. The mapping of DNAs to chromosomes according to the present invention is an important first step in correlating those sequences with genes associated with disease.

10 In certain preferred embodiments in this regard, the cDNA herein disclosed is used to clone genomic DNA of a NKX3.1 protein gene. This can be accomplished using a variety of well known techniques and libraries, which generally are available commercially. The genomic DNA then is used for *in situ* chromosome mapping using well known techniques for this purpose.

15 In addition, in some cases, sequences can be mapped to chromosomes by preparing PCR primers (preferably 15-25 bp) from the cDNA. Computer analysis of the 3' untranslated region of the gene is used to rapidly select primers that do not span more than one exon in the genomic DNA, thus complicating the amplification process. These primers are then used for PCR screening of somatic cell hybrids containing individual human chromosomes.

20 Fluorescence *in situ* hybridization ("FISH") of a cDNA clone to a metaphase chromosomal spread can be used to provide a precise chromosomal location in one step. This technique can be used with probes from the cDNA as short as 50 or 60 bp. For a review of this technique, see Verma *et al.*, *Human Chromosomes: A Manual Of Basic Techniques*, Pergamon Press, New York
25 (1988).

Once a sequence has been mapped to a precise chromosomal location, the physical position of the sequence on the chromosome can be correlated with

genetic map data. Such data are found, for example, in V. McKusick, *Mendelian Inheritance In Man*, available on-line through Johns Hopkins University, Welch Medical Library. The relationship between genes and diseases that have been mapped to the same chromosomal region are then identified through linkage analysis (coinheritance of physically adjacent genes).

Next, it is necessary to determine the differences in the cDNA or genomic sequence between affected and unaffected individuals. If a mutation is observed in some or all of the affected individuals but not in any normal individuals, then the mutation is likely to be the causative agent of the disease.

The NKX3.1 gene appears to be located on 8p21, a region which is deleted in up to 60-80% of prostate cancers and which suggests a role for the gene in androgen-driven maintenance of prostate tissue phenotype. (See, Example 6 below).

Having generally described the invention, the same will be more readily understood by reference to the following examples, which are provided by way of illustration and are not intended as limiting.

Examples

Example 1: Expression and Purification of NKX3.1 in E. coli

The bacterial expression vector pQE9 (pD10) is used for bacterial expression in this example. (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311). pQE9 encodes ampicillin antibiotic resistance ("Amp^r") and contains a bacterial origin of replication ("ori"), an IPTG inducible promoter, a ribosome binding site ("RBS"), six codons encoding histidine residues that allow affinity purification using nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin sold by QIAGEN, Inc., *supra*, and suitable single restriction enzyme cleavage sites. These elements are arranged such that an inserted DNA fragment encoding a polypeptide expresses that polypeptide with the six His residues (i.e., a "6 X His tag")) covalently linked to the amino terminus of that polypeptide.

The DNA sequence encoding the desired portion NKX3.1 protein lacking the hydrophobic leader sequence is amplified from the deposited cDNA clone using PCR oligonucleotide primers which anneal to the amino terminal sequences of the desired portion of the NKX3.1 protein and to sequences in the deposited construct 3' to the cDNA coding sequence. Additional nucleotides containing restriction sites to facilitate cloning in the pQE9 vector are added to the 5' and 3' primer sequences, respectively.

For cloning the mature protein, the 5' primer has the sequence 5' GCGGGATCCATGCTCAGGGTTCGGAG 3' (SEQ ID NO: 15) containing the underlined BamHI restriction site followed by 18 nucleotides complementary to the amino terminal coding sequence of the NKX3.1 sequence in Figure 1. One of ordinary skill in the art would appreciate, of course, that the point in the protein coding sequence where the 5' primer begins may be varied to amplify a DNA segment encoding any desired portion of the complete NKX3.1 protein shorter or longer than the complete sequence. The 3' primer has the sequence 5' GCGAGCTTTTACCCAAAAGCTGGGCT 3' (SEQ ID NO:16) containing the underlined HindIII restriction site followed by 18 nucleotides complementary to the non-coding sequence of the NKX3.1 DNA sequence in Figure 1.

The amplified NKX3.1 DNA fragment and the vector pQE9 are digested with BamHI and HindIII and the digested DNAs are then ligated together. Insertion of the NKX3.1 DNA into the restricted pQE9 vector places the NKX3.1 protein coding region downstream from the IPTG-inducible promoter and in-frame with an initiating AUG and the six histidine codons.

The ligation mixture is transformed into competent *E. coli* cells using standard procedures such as those described in Sambrook *et al.*, *Molecular Cloning: a Laboratory Manual, 2nd Ed.*; Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY (1989). *E. coli* strain M15/rep4, containing multiple copies of the plasmid pREP4, which expresses the lac repressor and confers kanamycin resistance ("Kan"), is used in carrying out the illustrative example described herein. This strain, which is only one of many that are suitable for

expressing NKX3.1 protein, is available commercially from QIAGEN, Inc., *supra*. Transformants are identified by their ability to grow on LB plates in the presence of ampicillin and kanamycin. Plasmid DNA is isolated from resistant colonies and the identity of the cloned DNA confirmed by restriction analysis, PCR and DNA sequencing.

Clones containing the desired constructs are grown overnight ("O/N") in liquid culture in LB media supplemented with both ampicillin (100 µg/ml) and kanamycin (25 µg/ml). The O/N culture is used to inoculate a large culture, at a dilution of approximately 1:25 to 1:250. The cells are grown to an optical density at 600 nm ("OD600") of between 0.4 and 0.6. Isopropyl-b-D-thiogalactopyranoside ("IPTG") is then added to a final concentration of 1 mM to induce transcription from the lac repressor sensitive promoter, by inactivating the lacI repressor. Cells subsequently are incubated further for 3 to 4 hours. Cells then are harvested by centrifugation.

The cells are then stirred for 3-4 hours at 4 °C in 6M guanidine-HCl, pH8. The cell debris is removed by centrifugation, and the supernatant containing the NKX3.1 is loaded onto a nickel-nitrilo-tri-acetic acid ("NiNTA") affinity resin column (available from QIAGEN, Inc., *supra*). Proteins with a 6 x His tag bind to the Ni-NTA resin with high affinity and can be purified in a simple one-step procedure (for details see: The QIAexpressionist, 1995, QIAGEN, Inc., *supra*). Briefly the supernatant is loaded onto the column in 6 M guanidine-HCl, pH8, the column is first washed with 10 volumes of 6 M guanidine-HCl, pH8, then washed with 10 volumes of 6 M guanidine-HCl pH6, and finally the NKX3.1 is eluted with 6 M guanidine-HCl, pH5.

The purified protein is then renatured by dialyzing it against phosphate-buffered saline (PBS) or 50 mM Na-acetate, pH 6 buffer plus 200 mM NaCl. Alternatively, the protein can be successfully refolded while immobilized on the Ni-NTA column. The recommended conditions are as follows: renature using a linear 6M-1M urea gradient in 500 mM NaCl, 20% glycerol, 20 mM Tris/HCl pH7.4, containing protease inhibitors. The renaturation should be performed over

a period of 1.5 hours or more. After renaturation the proteins can be eluted by the addition of 250 mM imidazole. Imidazole is removed by a final dialyzing step against PBS or 50 mM sodium acetate pH6 buffer plus 200 mM NaCl. The purified protein is stored at 4°C or frozen at -80°C.

5 ***Example 2: Cloning and Expression of NKX3.1 protein in a Baculovirus Expression System***

10 In this illustrative example, the plasmid shuttle vector pA2 GP is used to insert the cloned DNA encoding the protein into a baculovirus to express the NKX3.1 protein, using a baculovirus leader and standard methods as described in Summers *et al.*, *A Manual of Methods for Baculovirus Vectors and Insect Cell Culture Procedures*, Texas Agricultural Experimental Station Bulletin No. 1555 (1987). This expression vector contains the strong polyhedrin promoter of the *Autographa californica* nuclear polyhedrosis virus (AcMNPV) followed by the secretory signal peptide (leader) of the baculovirus gp67 protein and convenient
15 restriction sites such as BamHI, Xba I and Asp718. The polyadenylation site of the simian virus 40 ("SV40") is used for efficient polyadenylation. For easy selection of recombinant virus, the plasmid contains the beta-galactosidase gene from *E. coli* under control of a weak *Drosophila* promoter in the same orientation, followed by the polyadenylation signal of the polyhedrin gene. The inserted genes
20 are flanked on both sides by viral sequences for cell-mediated homologous recombination with wild-type viral DNA to generate viable virus that expresses the cloned polynucleotide.

25 Many other baculovirus vectors could be used in place of the vector above, such as pAc373, pVL941 and pAcIM1, as one skilled in the art would readily appreciate, as long as the construct provides appropriately located signals for transcription, translation, secretion and the like, including a signal peptide and an in-frame AUG as required. Such vectors are described, for instance, in Luckow *et al.*, *Virology* 170:31-39.

The cDNA sequence encoding the NKX3.1 protein in the deposited clone shown in Figure 1 (SEQ ID NO:2), is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' sequences of the gene.

The 5' primer has the sequence 5' GCGGGATCCCATGCTCAGGGTCCGGAG 3' (SEQ ID NO:17) containing the underlined BamHI restriction enzyme site followed by 18 bases of the sequence of the mature NKX3.1 protein shown in Figure 1, beginning with the indicated N-terminus of the mature protein. The 3' primer has the sequence 5'GCGGATCCTTACCCAAAAGCTGGGCT 3'(SEQ ID NO:18) containing the underlined BamHI restriction site followed by 18 nucleotides complementary to the 3' noncoding sequence in Figure 1.

The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("Geneclean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with BamHI and again is purified on a 1% agarose gel. This fragment is designated herein "F1".

The plasmid is digested with the restriction enzymes BamHI and optionally, can be dephosphorylated using calf intestinal phosphatase, using routine procedures known in the art. The DNA is then isolated from a 1% agarose gel using a commercially available kit ("Geneclean" BIO 101 Inc., La Jolla, Ca.). This vector DNA is designated herein "V1".

Fragment F1 and the dephosphorylated plasmid V1 are ligated together with T4 DNA ligase. *E. coli* HB101 or other suitable *E. coli* hosts such as XL-1 Blue (Stratagene Cloning Systems, La Jolla, CA) cells are transformed with the ligation mixture and spread on culture plates. Bacteria are identified that contain the plasmid with the human NKX3.1 gene using the PCR method, in which one of the primers that is used to amplify the gene and the second primer is from well within the vector so that only those bacterial colonies containing the NKX3.1 gene fragment will show amplification of the DNA. The sequence of the cloned fragment is confirmed by DNA sequencing. This plasmid is designated herein pBacNKX3.1.

Five μg of the plasmid pBacNKX3.1 is co-transfected with 1.0 μg of a commercially available linearized baculovirus DNA ("BaculoGold™ baculovirus DNA", Pharmingen, San Diego, CA.), using the lipofection method described by Felgner *et al.*, *Proc. Natl. Acad. Sci. USA* 84:7413-7417 (1987). 1 μg of BaculoGold™ virus DNA and 5 μg of the plasmid pBacNKX3.1 are mixed in a sterile well of a microtiter plate containing 50 μl of serum-free Grace's medium (Life Technologies Inc., Gaithersburg, MD). Afterwards, 10 μl Lipofectin plus 90 μl Grace's medium are added, mixed and incubated for 15 minutes at room temperature. Then the transfection mixture is added drop-wise to Sf9 insect cells (ATCC CRL 1711) seeded in a 35 mm tissue culture plate with 1 ml Grace's medium without serum. The plate is rocked back and forth to mix the newly added solution. The plate is then incubated for 5 hours at 27°C. After 5 hours the transfection solution is removed from the plate and 1 ml of Grace's insect medium supplemented with 10% fetal calf serum is added. The plate is put back into an incubator and cultivation is continued at 27°C for four days.

After four days the supernatant is collected and a plaque assay is performed, as described by Summers and Smith, *supra*. An agarose gel with "Blue Gal" (Life Technologies Inc., Gaithersburg) is used to allow easy identification and isolation of gal-expressing clones, which produce blue-stained plaques. (A detailed description of a "plaque assay" of this type can also be found in the user's guide for insect cell culture and baculovirology distributed by Life Technologies Inc., Gaithersburg, page 9-10). After appropriate incubation, blue stained plaques are picked with the tip of a micropipettor (e.g., Eppendorf). The agar containing the recombinant viruses is then resuspended in a microcentrifuge tube containing 200 μl of Grace's medium and the suspension containing the recombinant baculovirus is used to infect Sf9 cells seeded in 35 mm dishes. Four days later the supernatants of these culture dishes are harvested and then they are stored at 4°C. The recombinant virus is called V-NKX3.1.

To verify the expression of the NKX3.1 gene, Sf9 cells are grown in Grace's medium supplemented with 10% heat inactivated FBS. The cells are

infected with the recombinant baculovirus V-NKX3.1 at a multiplicity of infection ("MOI") of about 2. Six hours later the medium is removed and is replaced with SF900 II medium minus methionine and cysteine (available from Life Technologies Inc., Rockville, MD). If radiolabeled proteins are desired, 42 hours later, 5 μ Ci of 35 S-methionine and 5 μ Ci 35 S-cysteine (available from Amersham) are added. The cells are further incubated for 16 hours and then they are harvested by centrifugation. The proteins in the supernatant as well as the intracellular proteins are analyzed by SDS-PAGE followed by autoradiography (if radiolabeled). Microsequencing of the amino acid sequence of the amino terminus of purified protein may be used to determine the amino terminal sequence of the mature protein and thus the cleavage point and length of the secretory signal peptide.

Example 3: Cloning and Expression of NKX3.1 in Mammalian Cells

A typical mammalian expression vector contains the promoter element, which mediates the initiation of transcription of mRNA, the protein coding sequence, and signals required for the termination of transcription and polyadenylation of the transcript. Additional elements include enhancers, Kozak sequences and intervening sequences flanked by donor and acceptor sites for RNA splicing. Highly efficient transcription can be achieved with the early and late promoters from SV40, the long terminal repeats (LTRS) from Retroviruses, e.g., RSV, HTLVI, HIVI and the early promoter of the cytomegalovirus (CMV). However, cellular elements can also be used (e.g., the human actin promoter). Suitable expression vectors for use in practicing the present invention include, for example, vectors such as PSVL and PMSG (Pharmacia, Uppsala, Sweden), pRSVcat (ATCC 37152), pSV2dhfr (ATCC 37146) and pBC12MI (ATCC 67109). Mammalian host cells that could be used include, human Hela 293, H9 and Jurkat cells, mouse NIH3T3 and C127 cells, Cos 1, Cos 7 and CV 1, quail QC1-3 cells, mouse L cells and Chinese hamster ovary (CHO) cells.

Alternatively, the gene can be expressed in stable cell lines that contain the gene integrated into a chromosome. The co-transfection with a selectable marker such as dhfr, gpt, neomycin, or hygromycin allows the identification and isolation of the transfected cells.

5 The transfected gene can also be amplified to express large amounts of the encoded protein. The DHFR (dihydrofolate reductase) marker is useful to develop cell lines that carry several hundred or even several thousand copies of the gene of interest. Another useful selection marker is the enzyme glutamine synthase (GS) (Murphy *et al.*, *Biochem J.* 227:277-279 (1991); Bebbington *et al.*,
10 *Bio/Technology* 10:169-175 (1992)). Using these markers, the mammalian cells are grown in selective medium and the cells with the highest resistance are selected. These cell lines contain the amplified gene(s) integrated into a chromosome. Chinese hamster ovary (CHO) and NSO cells are often used for the production of proteins.

15 The expression vectors pC1 and pC4 contain the strong promoter (LTR) of the Rous Sarcoma Virus (Cullen *et al.*, *Molecular and Cellular Biology*, 438-447 (March, 1985)) plus a fragment of the CMV-enhancer (Boshart *et al.*, *Cell* 41:521-530 (1985)). Multiple cloning sites, e.g., with the restriction enzyme cleavage sites BamHI, XbaI and Asp718, facilitate the cloning of the gene of
20 interest. The vectors contain in addition the 3' intron, the polyadenylation and termination signal of the rat preproinsulin gene.

Example 3A: Cloning and Expression in COS Cells

25 The expression plasmid, pNKX3.1 HA, is made by cloning a cDNA encoding NKX3.1 into the expression vector pcDNAI/Amp or pcDNAIII (which can be obtained from Invitrogen, Inc.).

 The expression vector pcDNAI/amp contains: (1) an *E. coli* origin of replication effective for propagation in *E. coli* and other prokaryotic cells; (2) an ampicillin resistance gene for selection of plasmid-containing prokaryotic cells; (3)

an SV40 origin of replication for propagation in eukaryotic cells; (4) a CMV promoter, a polylinker, an SV40 intron; (5) several codons encoding a hemagglutinin fragment (i.e., an "HA" tag to facilitate purification) followed by a termination codon and polyadenylation signal arranged so that a cDNA can be conveniently placed under expression control of the CMV promoter and operably linked to the SV40 intron and the polyadenylation signal by means of restriction sites in the polylinker. The HA tag corresponds to an epitope derived from the influenza hemagglutinin protein described by Wilson *et al.*, *Cell* 37:767 (1984). The fusion of the HA tag to the target protein allows easy detection and recovery of the recombinant protein with an antibody that recognizes the HA epitope. pcDNAIII contains, in addition, the selectable neomycin marker.

A DNA fragment encoding the NKX3.1 is cloned into the polylinker region of the vector so that recombinant protein expression is directed by the CMV promoter. The plasmid construction strategy is as follows. The NKX3.1 cDNA of the deposited clone is amplified using primers that contain convenient restriction sites, much as described above for construction of vectors for expression of NKX3.1 in *E. coli*. Suitable primers include the following, which are used in this example. The 5' primer, containing the underlined BamHI site, a Kozak sequence, an AUG start codon and 18 bases of the 5' coding region of the complete NKX3.1 has the following sequence: 5'GCGGGATCCCATGCTCAGGGTTCCGGAG 3' (SEQ ID NO:17). The 3' primer, containing the underlined BamHI site, a stop codon, and 18 bases of 3' coding sequence has the following sequence (at the 3' end): 5'GCGGATCCTTACCCAAAAGCTGGGCT 3' (SEQ ID NO:18).

The PCR amplified DNA fragment and the vector, pcDNAI/Amp, are digested with BamHI and then ligated. The ligation mixture is transformed into *E. coli* strain SURE (available from Stratagene Cloning Systems, 11099 North Torrey Pines Road, La Jolla, CA 92037), and the transformed culture is plated on ampicillin media plates which then are incubated to allow growth of ampicillin resistant colonies. Plasmid DNA is isolated from resistant colonies and examined

by restriction analysis or other means for the presence of the NKX3.1-encoding fragment.

For expression of recombinant NKX3.1, COS cells are transfected with an expression vector, as described above, using DEAE-DEXTRAN, as described, for instance, in Sambrook *et al.*, *Molecular Cloning: a Laboratory Manual*, Cold Spring Laboratory Press, Cold Spring Harbor, New York (1989). Cells are incubated under conditions for expression of NKX3.1 by the vector.

Expression of the NKX3.1-HA fusion protein is detected by radiolabeling and immunoprecipitation, using methods described in, for example Harlow *et al.*, *Antibodies: A Laboratory Manual, 2nd Ed.*; Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York (1988). To this end, two days after transfection, the cells are labeled by incubation in media containing ³⁵S-cysteine for 8 hours. The cells and the media are collected, and the cells are washed and lysed with detergent-containing RIPA buffer: 150 mM NaCl, 1% NP-40, 0.1% SDS, 0.5% DOC, 50 mM TRIS, pH 7.5, as described by Wilson *et al.* cited above. Proteins are precipitated from the cell lysate and from the culture media using an HA-specific monoclonal antibody. The precipitated proteins then are analyzed by SDS-PAGE and autoradiography. An expression product of the expected size is seen in the cell lysate, which is not seen in negative controls.

Example 3B: Cloning and Expression in CHO Cells

The vector pC4 is used for the expression of NKX3.1 protein. Plasmid pC4 is a derivative of the plasmid pSV2-dhfr (ATCC Accession No. 37146). The plasmid contains the mouse DHFR gene under control of the SV40 early promoter. Chinese hamster ovary- or other cells lacking dihydrofolate activity that are transfected with these plasmids can be selected by growing the cells in a selective medium (alpha minus MEM, Life Technologies) supplemented with the chemotherapeutic agent methotrexate. The amplification of the DHFR genes in cells resistant to methotrexate (MTX) has been well documented (see, e.g., Alt,

F. W., Kellems, R. M., Bertino, J. R., and Schimke, R. T., 1978, *J Biol. Chem.* 253:1357-1370, Hamlin, J. L. and Ma, C. 1990, *Biochem. et Biophys. Acta*, 1097:107-143, Page, M. J. and Sydenham, M.A. 1991, *Biotechnology* 9:64-68).

Cells grown in increasing concentrations of MTX develop resistance to the drug by overproducing the target enzyme, DHFR, as a result of amplification of the DHFR gene. If a second gene is linked to the DHFR gene, it is usually co-amplified and over-expressed. It is known in the art that this approach may be used to develop cell lines carrying more than 1,000 copies of the amplified gene(s). Subsequently, when the methotrexate is withdrawn, cell lines are obtained which contain the amplified gene integrated into one or more chromosome(s) of the host cell.

Plasmid pC4 contains for expressing the gene of interest the strong promoter of the long terminal repeat (LTR) of the Rous Sarcoma Virus (Cullen, *et al.*, *Molecular and Cellular Biology*, March 1985:438-447) plus a fragment isolated from the enhancer of the immediate early gene of human cytomegalovirus (CMV) (Boshart *et al.*, *Cell* 41:521-530 (1985)). Downstream of the promoter are BamHI, XbaI, and Asp718 restriction enzyme cleavage sites that allow integration of the genes. Behind these cloning sites the plasmid contains the 3' intron and polyadenylation site of the rat preproinsulin gene. Other high efficiency promoters can also be used for the expression, e.g., the human β -actin promoter, the SV40 early or late promoters or the long terminal repeats from other retroviruses, e.g., HIV and HTLV. Clontech's Tet-Off and Tet-On gene expression systems and similar systems can be used to express the NKX3.1 in a regulated way in mammalian cells (Gossen, M., & Bujard, H. 1992, *Proc. Natl. Acad. Sci. USA* 89: 5547-5551). For the polyadenylation of the mRNA other signals, e.g., from the human growth hormone or globin genes can be used as well. Stable cell lines carrying a gene of interest integrated into the chromosomes can also be selected upon co-transfection with a selectable marker such as gpt, G418 or hygromycin. It is advantageous to use more than one selectable marker in the beginning, e.g., G418 plus methotrexate.

The plasmid pC4 is digested with the restriction enzymes BamHI and then dephosphorylated using calf intestinal phosphatase by procedures known in the art. The vector is then isolated from a 1% agarose gel.

The DNA sequence encoding the complete NKX3.1 protein including its leader sequence is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' sequences of the gene. The 5' primer has the sequence 5'GCGGGATCCCATGCTCAGGGTTCGGAG 3' (SEQ ID NO:17) containing the underlined BamHI restriction enzyme site followed by an efficient signal for initiation of translation in eukaryotes, as described by Kozak, M., *J. Mol. Biol.* 196:947-950 (1987), and 18 bases of the coding sequence of NKX3.1 shown in Figure 1 (SEQ ID NO:1). The 3' primer has the sequence 5' GCGGATCCTTACCCAAAAGCTGGGCT 3' (SEQ ID NO:19) containing the underlined BamHI restriction site followed by 18 nucleotides complementary to the non-translated region of the NKX3.1 gene shown in Figure 1 (SEQ ID NO:1).

The amplified fragment is digested with the endonuclease BamHI and then purified again on a 1% agarose gel. The isolated fragment and the dephosphorylated vector are then ligated with T4 DNA ligase. *E. coli* HB101 or XL-1 Blue cells are then transformed and bacteria are identified that contain the fragment inserted into plasmid pC4 using, for instance, restriction enzyme analysis.

Chinese hamster ovary cells lacking an active DHFR gene are used for transfection. 5 µg of the expression plasmid pC4 is cotransfected with 0.5 µg of the plasmid pSV2-neo using lipofectin (Felgner *et al.*, *supra*). The plasmid pSV2-neo contains a dominant selectable marker, the neo gene from Tn5 encoding an enzyme that confers resistance to a group of antibiotics including G418. The cells are seeded in alpha minus MEM supplemented with 1 mg/ml G418. After 2 days, the cells are trypsinized and seeded in hybridoma cloning plates (Greiner, Germany) in alpha minus MEM supplemented with 10, 25, or 50 ng/ml of methotrexate plus 1 mg/ml G418. After about 10-14 days single clones are trypsinized and then seeded in 6-well petri dishes or 10 ml flasks using different concentrations of methotrexate (50 nM, 100 nM, 200 nM, 400 nM, 800 nM).

Clones growing at the highest concentrations of methotrexate are then transferred to new 6-well plates containing even higher concentrations of methotrexate (1 μ M, 2 μ M, 5 μ M, 10 mM, 20 mM). The same procedure is repeated until clones are obtained which grow at a concentration of 100 - 200 μ M. Expression of the desired gene product is analyzed, for instance, by SDS-PAGE and Western blot or by reverse phase HPLC analysis.

Example 4: Tissue distribution of NKX3.1 mRNA expression

To examine the tissue specificity of NKX3.1 expression, two separate Northern analyses were performed using mRNA derived from overlapping sets of human tissues using methods described by, among others, Sambrook *et al.*, cited above.

Materials and Methods

A cDNA probe containing the entire nucleotide sequence of the NKX3.1 protein (SEQ ID NO: 1) was labeled with 32 P using the *rediprime*TM DNA labeling system (Amersham Life Science), according to manufacturer's instructions. After labeling, the probe was purified using a CHROMA SPIN-100TM column (Clontech Laboratories, Inc.), according to manufacturer's protocol number PT1200-1. The purified labeled probe was then used to examine various human tissues for NKX3.1 mRNA.

Multiple Tissue Northern (MTN) blots containing various human tissues (H) or human immune system tissues (IM) are obtained from Clontech and are examined with the labeled probe using ExpressHybTM hybridization solution (Clontech) according to manufacturer's protocol number PT1190-1. Following hybridization and washing, the blots are mounted and exposed to film at -70°C overnight, and films developed according to standard procedures.

Cell lines were grown in medium containing 10% fetal bovine serum. Human peripheral blood lymphocytes (HuPBL) were isolated from a normal

donor (Ficoll-Paque, Pharmacia). Total cellular RNA was isolated from each of the cell lines using the Trizol RNA isolation reagent (Gibco BRL). Ten microgram aliquots were resolved on a 1% agarose-formaldehyde gels and transferred to a nylon membrane (Hybond-N, Amersham) essentially as described (Sambrook *et al.*, 1988). Filters were prehybridized for 1 hour in SLURP (7 mM Tris-HCl pH 7.5, 4x SSC, 10% dextran sulfate, 0.8x Denhardts solution, 40% formamide, 20 mg/mL salmon sperm DNA and 0.5% SDS). NKX3.1 or glyceraldehyde-3-phosphate dehydrogenase (GAPDH) cDNAs were labeled with ³²P dCTP by random priming, denatured, added to the prehybridization mix, and hybridized overnight at 42°C. Filters were washed at 42°C for 10 min, then at room temperature, in 2x SSC/0.1%SDS, followed by a high stringency wash at 65°C for 40 min in 0.2x SSC/0.1%SDS. Washed filters were then dried and exposed to X-ray film.

Results

An abundant 3.5kb NKX3.1 mRNA was detected in prostate and at much lower level in testis. NKX3.1 mRNA was not detected in several other adult tissues including brain, kidney, small intestine, pancreas, heart, liver, lung, thymus, spleen, placenta, colon, lymphocytes and ovary. We also examined the expression of NKX3.1 mRNA in a variety of cultured cells including both transformed and non-transformed human cell lines (Table 1, below). NKX3.1 expression was seen only in the hormone-responsive, androgen receptor-positive LNCaP prostate cancer cell line. However NKX3.1 expression was not observed in either of the two androgen receptor-negative cell lines, PC-3 and DU-145, nor in eleven other cell lines of varied tissue origin. Interestingly, other HOX genes tested were expressed in several of the same cell-lines indicating even more strongly that NKX3.1 exhibits a uniquely restricted tissue expression pattern. These findings indicate that NKX3.1 is indeed a new prostate-specific gene. Since NKX3.1 is a member of the homeobox family, it may be involved in establishing the developmental cascade of specific cell types, suggesting that this gene may have

a role in the development and differentiation of prostatic tissue. It is also intriguing that the NKX3.1 expression has apparently been lost in both of the androgen independent cell lines tested, but not in the androgen-dependent cell line LNCaP, indicating that expression of the gene may be androgen-regulated.

5

Table 1

Cell lines used for Northern blot analysis
Cell Line (Tissue)

10

FS4 (Human diploid fibroblast) ¹	RT-4 (Bladder Carcinoma) ⁵
HuPBL (Peripheral Blood Lymphocyte)	HTB-44 (Kidney Carcinoma) ⁵
DAOY (Medulloblastoma) ²	OVCAR-3 (Ovarian Carcinoma) ⁵
BHM22 (B cell Leukemia) ³	CATES-1B (Embryonal Carcinoma) ²
8392 (EBV-Transformed B-cell) ³	PC3 (Prostate Carcinoma) ⁶
HELA (Cervical Carcinoma) ⁴	LNCaP (Prostate Carcinoma) ⁶
SW480 (Colon Carcinoma) ³	DU-145 (Prostate Carcinoma) ²

15

Cell lines were obtained from the following sources and/or generously provided by the following laboratories:

¹ J. Vilcek, NYU Medical Center; ² ATCC; ³ A. Rabson, Center for advanced Biotechnology and Medicine; ⁴ C. Gelinas, Center for Advanced Biotechnology and Medicine; ⁵ The Cancer Institute of New Jersey; ⁶ S. Ward, University of Medicine and Dentistry New Jersey.

20

Example 5A: Androgen regulation of NKX3.1

25

Since NKX3.1 was expressed in LNCaP cells, which are dependent on the presence of physiologic levels of androgen for growth and tumorigenesis (Burnes *et al.*, *Prostate* 9:247-259 (1986); Olea *et al.*, *Endocrinology* 126:1457-1463 (1990)), we examined whether NKX3.1 mRNA levels are responsive to androgen stimulation.

Materials and Methods

For the androgen regulation experiments, parallel cultures of PC3 or LNCaP cells were incubated in fetal bovine serum (10%) containing medium or were incubated in medium containing 10% charcoal dextran treated human serum (Sigma) as indicated. The synthetic androgen R1881 (NEN-Dupont) was dissolved in 100% ethanol. Mock induced cultures were treated with an equal volume of 100% ethanol alone. Northern analysis was then performed as described above in Example 4.

Results

In standard serum-containing growth medium (control cells), NKX3.1 mRNA was detected in Northern analyses at a level similar to that seen in the previous cell-line survey. However, when cells were grown in the absence of androgen stimulation, NKX3.1 message was reduced to undetectable levels. Addition of the synthetic androgen R1881 to these cultures in concentrations as low as 0.3 nM resulted in an restoration NKX3.1 mRNA to control levels and higher concentrations of R1881 resulted in a dose-dependent increase in the expression gene. In contrast, NKX3.1 mRNA was not detected in the androgen-independent prostate carcinoma line PC-3 in either the absence or presence of R1881. Experiments have demonstrated that NKX3.1 expression in LNCaP cells is exquisitely sensitive to the presence or absence of androgens and that androgen stimulation is at the transcription level and does not require synthesis of additional cellular proteins upon stimulation of the hormone. These results raise the possibility that NKX3.1 is part of the prostate cell's primary response to androgen stimulation and indicate that the gene is a candidate for playing a central role in the differentiation of normal prostatic tissue and the reversal of normal differentiation seen during cancer progression.

Example 5B: NKX3.1 regulation in response to orchidectomy

The maintenance of differentiated functions within the prostate is well-established to be androgen-dependent (Davis, P. and Eaton, C.L., *J. Endocrinol.* 131:5-17 (1991)). Castration-induced androgen deprivation leads to a rapid shut-off of genes encoding prostate-specific secretory proteins (Mills, L.S. *et al.*, *EMBO J.* 6:3711-3717 (1987)). To determine whether *Nkx-3.1* was regulated in response to orchidectomy, RNA was extracted from prostates harvested at various time-points after castration, but prior to the onset of an atrophic state.

Materials and Methods

Orchidectomy was performed on 6-week old CD-1 mice as described for rats (Waynforth, H.B., *Experimental and Surgical Techniques in the Rat*, Academic Press Inc., San Diego, CA (1980)). RNA was extracted from total prostate, pooled from two mice, at each time-point after castration. Densitometric analysis of Northern blot autoradiograms was performed using Bioimage Software version 4.6P (Bioimage Inc., Ann Arbor, MI).

Results

Northern blot analysis of the RNA extracted from prostates revealed that by 24 hours after castration, the steady-state level of NKX3.1 mRNA was decreased nearly 10-fold. By 96 hours, the level was decreased 30-fold. These data suggest that the maintenance of a high level of expression of NKX3.1 requires testicular androgens. The same RNA blot was subsequently hybridized with a probe that detected the mRNA encoding a secreted protease inhibitor, *mp12*, that has been demonstrated to be androgen-dependent (Mills, L.S. *et al.*, *EMBO J.* 6:3711-3717 (1987)). A comparison of the kinetics and extent of downregulation showed that the level of *mp12* mRNA was decreased by more than 70-fold by 24 hours after castration. At 96 hours, *mp12* mRNA was no longer detectable by Northern analysis, whereas NKX3.1 mRNA fell to a basal

level that was maintained for at least several more days. These data suggest that NKX3.1 expression is androgen-responsive, but also show a low, basal level of expression that may not be androgen-dependent.

Example 6: Chromosomal Mapping of NKX3.1

5 To demonstrate the chromosomal location of the NKX3.1 gene, a 20 kb genomic NKX3.1 clone was used as a probe for fluorescence *in situ* hybridization to human chromosome metaphase spreads (Lawrence *et al.*, *Cell* 52:51-61 (1988)).

Materials and Methods

10 An NKX3-1 lambda genomic clone was isolated by standard techniques (Sambrook *et al.*, 1989). This DNA was nick-translated using digoxigenin-11-dUTP (Boehringer Mannheim) and *in situ* hybridization was done as detailed in Johnson *et al.*, *Methods in Cell Biol.* 35:73-99 (1991). Individual chromosomes were counterstained with DAPI and color digital images, containing both DAPI
15 and gene signal were recorded using a triple-band pass filter set (Chroma Technology, Inc. Brattleboro, VT) in combination with a charged couple-device camera (Photometrics, Inc. Tucson, AZ) and variable excitation wave length filters, which allows for recording multi-color images without image shift (Johnson
20 *et al.*, *Genetic analysis: Techniques and applications* 8:75-76 (1991)). In experiments analyzing the position of more than one gene or marker on 8p, each DNA probe was labeled separately with either digoxigenin or biotin (Gibco/BRL) and the two were mixed at different ratios for each DNA probe during hybridization in the triple-label experiments. The hybridized signal was subsequently detected using Rhodamine-conjugated anti-digoxigenin antibodies
25 and FITC-streptavidin (Boehringer-Mannheim). Individual probes were identified based on the red:green ratio of the recorded signal. Images were analyzed using

the ISEE software package (Inovision Corp. Durham, NC) and in some cases the gene signal was pseudo-colored in the final image for clarity.

Results

Approximately 20 metaphase chromosome spreads were analyzed by eye,
5 most of which had a doublet signal characteristic of genuine hybridization on at least one chromosome 8. Doublet signal was not detected on any other chromosome. Detailed analysis of 10 individual chromosomes, using a combination of fractional length measurements and fluorescence banding, combined with high-resolution image analysis, indicated that the NKX3.1 gene is
10 positioned within band 8p21. Allelic deletions associated with this region of 8p are frequently reported in prostate cancer tissue and are thought to occur at two sites, one within band 8p22 and other within 8p21, making the precise assignment of the NKX3.1 band position of critical importance. Therefore, the position of the NKX3.1 was confirmed by co-mapping the gene with markers on 8p21 and
15 preliminary evidence indicates that the gene is within a no more than a few megabases of the 8p21 site that is frequently deleted in prostate cancer cells. Because a gene within this region of 8p21 is thought to be involved in the loss of androgen-driven differentiated prostatic phenotype it is reasonable to suggest that this might be that gene. This is consistent with a role for NKX3.1 in the
20 maintenance of prostate differentiation.

Example 7: NKX3.1 homeodomain exhibits NK-like DNA binding specificity

The human and mouse NKX3.1 cDNAs have a high degree of sequence similarity of its homeodomain with that of *Drosophila* NK-3. Particularly well conserved are residues in helix III and the N-terminal arm, which are known to
25 determine the DNA binding specificity of other homeodomains (Damante, G. and Di Lauro, R., *Proc. Natl. Acad. Sci. USA* 88:5388-5392 (1991); Ebu Isaac, V. *et al.*, *Biochem.* 34:7127-7134 (1995)), and to contact DNA directly (Gehring, W.J.

et al., *Cell* 78:211-223 (1994)). NKX3.1 homeodomain contains a tyrosine residue at position 54, which is the most distinguishing feature of NK homeodomains, and is largely responsible for the atypical DNA binding specificities of this class of homeodomains relative to the prototypic (e.g.,
5 Antennadia-like) homeodomains (Chen, C.Y. and Schwartz, R.J., *J. Biol. Chem.* 270:15626-15633 (1995); Damante, G. and Di Lauro, R., *Proc. Natl. Acad. Sci. USA* 88:5388-5392 (1991); Damante, G. et al., *Nucleic Acids Res.* 22:3075-3083 (1994); Guazzi, S. et al., *EMBO J.* 9:3631-3639 (1990); Harvey, R.P., *Dev. Biol.* 178:203-216 (1996)). In particular, NK-2 have been shown to interact
10 preferentially with DNA sites that contain a "CAAG" core, rather than the prototype "TAAT" core recognized by most other homeodomains (Chen, C.Y. and Schwartz, R.J., *J. Biol. Chem.* 270:15626-15633 (1995); Damante, G. and Di Lauro, R., *Proc. Natl. Acad. Sci. USA* 88:5388-5392 (1991)).

To investigate the DNA binding specificity of the murine NKX3.1
15 homeodomain, gel mobility shift analyses were performed using highly purified recombinant protein. Given its high content of proline and arginine residues (which are inefficiently translated in *E. coli*), it was predicted that the full length murine NKX3.1 would be poorly expressed in bacterial cells (Abate, C. et al., *Mol. Cell. Biol.* 11:3624-3632 (1991)). Therefore, the region containing the
20 murine homeodomain (NKX3.1HD) was expressed as a hexahistidine fusion polypeptide in *E. coli*, and obtained highly purified protein by nickel affinity chromatography. Human NKX3.1HD was also produced and purified, and as anticipated, these proteins behaved identically with respect to their expression, purification and DNA binding properties.

25 **Materials and Methods**

In vitro transcription/ translation was performed using the pBluescript-NKX3.1 plasmid as a template with the TnT coupled transcription/translation system (Promega) according to the protocol of the manufacturer.

To produce the NKX3.1 homeodomain (NKX3.1 HD) in *E. coli*, sequences encoding amino acids 122 to 188 were amplified by PCR using the primers that contained restriction sites for cloning into the *Bam*HI and *Hind*III sites of the bacterial expression plasmid, pQE-9 (Quiagen). Polypeptides were produced in *E. coli* as hexahistidine fusion proteins and purified by nickel affinity chromatography as previously described (Catron, K.M. *et al.*, *Mol. Cell. Biol.* 13;2354-2365 (1993)).

The DNA binding experiments were performed as follows. Briefly, increasing amounts of protein were incubated in binding buffer [10 mM Tris-HCl (pH .6), 50 mM NaCl, 5% glycerol, 5% sucrose, 0.2 mM EDTA, 7.5 mM MgCl₂, 0.5 mg/mL bovine serum albumin, 0.1% NP-40, 10 mM DTT, 500 µg of poly-didC] with 4 x 10⁴ cpm of ³²P-labeled, double stranded oligonucleotide for 20 min at room temperature. The sequences of the DNA binding sites were as follows (top strand shown):

15 5' CACTGCCCAGTCAAGTGTCTTGA 3' [C site, Guazzi, S. *et al.*, *EMBO J.* 9:3631-3639 (1990))] (SEQ ID NO:20)

5' CACTGCCCAGTCACGTGTTCGTGA 3' (SEQ ID NO:21)

5' AACTAATTGGAGGC 3' [site 6, (Catron, K.M. *et al.*, *Mol. Cell. Biol.* 13;2354-2365 (1993))] (SEQ ID NO:22)

20 5' AACTACTTGGAGGC 3' [site 6-19, (Catron, K.M. *et al.*, *Mol. Cell. Biol.* 13;2354-2365 (1993))] (SEQ ID NO:23)

5' CTCTAATGGCTTTTCTC 3' [site BS2, (Guazzi, S. *et al.*, *EMBO J.* 9:3631-3639 (1990))] (SEQ ID NO:24)

Results

25 The DNA binding specificity of NKX3.1HD was examined by gel mobility shift analysis using various DNA sites identified for NK-like and Antennapedia-like homeodomains. The DNA sites were as follows: (i) the consensus DNA site for the NKX2.1 homeodomain (TCAAGTG) (Guazzi, S. *et al.*, *EMBO J.* 9:3631-3639 (1990)) and a mutated version of this site that contains nucleotide

substitutions within the CAAG core (TCACGTG); and (ii) the consensus DNA site identified for an Antennapedia-like homeodomain (Msx1) (CTAATTG), a mutated version of this site that contains nucleotide substitutions within the TAAT core (CTACTTG), or one that contains nucleotide substitutions 3' of the TAAT core (CTAATGG) (Catron, K.M. *et al.*, *Mol. Cell. Biol.* 13:2354-2365 (1993); Guazzi, S. *et al.*, *EMBO J.* 9:3631-3639 (1990)). To compare the relative DNA binding affinity of NKX3.1HD for these DNA sites, gel mobility shift analysis was performed with varying concentrations of the purified protein, using equimolar amounts of each site. Of the various DNA sites tested, NKX3.1HD interacted preferentially with the site containing the "CAAG" core. NKX3.1HD also bound to DNA sites containing the "TAAT" core, albeit with lower apparent affinity. The specificity of the interaction was apparent from the lack of NKX3.1HD binding to DNA sites containing substitutions within the "CAAG" or "TAAT" cores. These results demonstrate that NKX3.1HD exhibits similar DNA binding specificity as NK-2 homeodomains; namely a preferential association with DNA sites containing a "CAAG" core and relatively low affinity interaction with DNA sites containing a "TAAT" core.

Example 8: Expression of murine NKX3.1 within the prostate

The mouse prostate is comprised of four paired components that are heterogeneous both in morphology and function (Frohman, M., *Meth. Enzymol.* 218:340-356 (1993)). To further characterize expression of NKX3.1 expression within the prostate, the ventral prostate, the dorsolateral prostate, and the coagulating gland (anterior prostate) were dissected from adult animals. Microdissection of prostate glands into component lobes was performed as described (Sugimura, Y. *et al.*, *Biol. Reprod.* 34:961-971 (1986)) The dorsolateral prostate was further subdivided into its dorsal and lateral components (Sugimura, Y. *et al.*, *Biol. Reprod.* 34:961-971 (1986)).

To determine whether NKX3.1 mRNA was differentially distributed among the four lobes, Northern blot analysis was performed on RNA extracted from pools of individual lobes. All four lobes expressed NKX3.1 at a similar steady-state level relative to the housekeeping gene β -actin, suggesting that this gene may be important for regulating functions that are shared between them.

To characterize the cellular distribution of NKX3.1 within the prostate gland, *in situ* hybridization to histological sections of all four components was performed. *In situ* hybridization was performed essentially as described (Bogarad, L.D. *et al.*, *Develop. Biol.* 133:537-549 (1989)), using a 530-bp Bgl II-Pst I fragment derived entirely from the mouse NKX3.1 3' noncoding region. The architecture of the adult prostate is relatively simple, consisting principally of a series of branching ducts lined with secretory epithelial cells (Sugimura, Y. *et al.*, *Biol. Reprod.* 34:961-971 (1986)). The ducts are wrapped by condensed stromal sleeves with sparse connective tissue interspersed between the ducts. Analysis of serial sections hybridized with antisense and control sense probes, showed signal only with the antisense probe, confirming the specificity of hybridization. Within sections of each lobe, hybridization signal was detected exclusively over the epithelial cells lining the ducts. Stromal cells did not show detectable expression of NKX3.1. There was no distinguishable difference in the intensity of the signal over epithelial cells derived from individual lobes. These observations demonstrate that NKX3.1 mRNA is confined to epithelial cells in the adult prostate and are consistent with a role for this gene in the maintenance of differentiated functions in the prostate gland.

These observation strongly implicate NKX3.1 in the differentiation of the urethral epithelium into the highly-specialized prostate epithelium. Given that the NKX3.1 protein is likely to function as a transcription factor and its expression is restricted to the prostate anlage, it seems likely that the NKX3.1 gene is playing a primary role in driving the differentiation of the prostate gland. The continued expression of this gene in all four component lobes in adult animals suggests that it may further be involved in the maintenance of this tissue.

Example 9: NKX3.1 Expression During Mouse Development

To gain insights into the potential roles of NKX3.1, its expression in embryonic and adult tissues was examined. First, RNase protection analysis was performed using tissues obtained from late-gestation embryos and adults. Several tissues from day 17.5 p.c. embryos were surveyed and were found low levels to have of NKX3.1 expression in the kidney and developing gonad, barely detectable levels in the thyroid, and no detectable levels of expression in several other tissues examined. Given NKX3.1 expression in the developing urogenital system, tissues obtained from sexually immature (2-4 weeks old) male, sexually mature (8-12 weeks old) male, and sexually immature and mature female mice were also surveyed. NKX3.1 expression was observed in the prostate, seminal vesicle, and testis of sexually immature and mature male mice. Moreover, significantly higher levels of NKX3.1 were detected in the prostate and seminal vesicles of the sexually mature, versus immature, mice. In contrast, NKX3.1 was not detectable in tissues from the immature or mature female urogenital system.

To further verify these observations, the expression of NKX3.1 was examined by Northern blot analysis, using tissues obtained from adult male mice (8-12 weeks old). A transcript of about 3.2 kB was detected in total RNA prepared from seminal vesicle or a pooled sample of bladder and prostate, but not from several other tissues. Together with the results of RNase protection analysis, these findings demonstrate that NKX3.1 exhibits a limited tissue distribution in the developing urogenital system, and that its expression is most elevated in sexually mature males.

Both the prostate and seminal vesicles, which express high levels of NKX3.1, require androgens for their growth and development (Cunha, G.R., *Cancer* 74:1030-1044 (1994); Cunha, G.R. *et al.*, *Endocrine Reviews* 8:338-362 (1987)). Since expression of NKX3.1 was essentially undetectable in the prostate of castrated mice, together with results showing the human NKX3.1 is androgen-

regulated in prostate carcinoma cells, these data suggest that androgen stimulation is required for maintenance of NKX3.1 expression.

Localized expression of NKX3.1 during murine embryogenesis

5 To further delineate the expression pattern of NKX3.1 during murine embryogenesis, *in situ* hybridization was performed on cryosections from mouse embryos at days 14.5 through 17.5 of gestation. Based on the RNase protection results, the analysis was focused on the development of the male urogenital system.

Materials and Methods

10 ***Isolation of RNA and analysis of expression:*** Tissues were obtained by dissection from male or female Swiss-Webster mice, or day 17.5 *p.c.* embryos and freshly frozen (-70°C), or were purchased from Pel-Freez Biologicals, Inc. Frozen tissues from sham operated or castrated mice, at 2 days post surgery, were obtained from Taconic Farms, Inc. RNA was prepared using Trizol RNA
15 isolation reagent (GIBCO-BRL) according to the protocol of the manufacturer. RNase protection analysis was performed essentially as described (Krieg, P.A. and Melton, D.A., *Meth. Enzymol.* 155:397-415 (1987)) using antisense or sense riboprobes prepared from various subcloned fragments of the NKX3.1 cDNA. The antisense riboprobe specific for the L32 ribosomal protein gene was described
20 in (Shen, M.M. and Leder, P., *Proc. Natl. Acad. Sci. USA* 89:8240-8244 (1992)). The RNase protection assays were performed using probe A which spans the intron-exon splice junction; similar results were obtained using probes B or C. Northern blot analysis was performed essentially as described (Ausubel, F. *et al.*, *In Current Protocols in Molecular Biology*, K. Janssen, Ed., John Wiley and
25 Sons, Inc., New York, New York (1995)) using a nylon filter membrane (Hybond-N, Amersham). Filters were prehybridized at 42°C for 1 hour in SLURP (7 mM Tris-HCl (pH 7.5), 4 x SSC, 10% dextran sulfate, 0.8x Denhardt's solution, 40% formamide, 20 mg/mL salmon sperm DNA and 0.5% SDS). Probes were labeled

with ^{32}P -dCTP by random priming (Ausubel, F. *et al.*, *In Current Protocols in Molecular Biology*, K. Janssen, Ed., John Wiley and Sons, Inc., New York, New York (1995)), and filters were hybridized overnight at 42°C. Filters were washed in 2x SSC/0.1% SDS at 42°C for 10 min, then at room temperature for 10 min, followed by a high stringency wash in 0.2x SSC/0.1% SDS at 65°C for 40 min.

In situ hybridization: Digoxigenin-labeled riboprobes, corresponding to regions of the Nkx3.1 cDNA (probes D, E, F) were synthesized as described (Catron, K.M. *et al.*, *Mech. Dev.* 55:185-199 (1996)). To determine the sex of Swiss Webster mouse embryos, genomic DNA was isolated from tails and PCR was performed using primers directed against the *Sry* gene (Hogan, B. *et al.*, *Manipulating the mouse embryo*, Cold Spring Harbor Laboratory Press, Cold Spring Harbor (1994)). Embryos were fixed with 4% paraformaldehyde/phosphate buffered saline (PBS) at 4°C for two days and transferred to a solution of 30% sucrose /PBS plus 0.1% Tween (PBT), bleached with 6% H_2O_2 / PBT, then washed three times with PBT. Sections were treated with 1 $\mu\text{g}/\text{ml}$ roteinase K/PBT, followed by sequential washes with 2 mg/ml glycine/PBT and PBT. Sections were incubated in prehybridization solution (50% formamide, 5X SSC (pH 4.5), 50 $\mu\text{g}/\text{ml}$ yeast tRNA, 1% SDS, 50 $\mu\text{g}/\text{ml}$ heparin) at 65°C for 1 hr. followed by hybridization with digoxigenin-labeled riboprobes overnight at 65°C. Sections were washed three times each in: 50% formamide/5X SSC (pH 4.5)/1% SDS at 70°C; 50% formamide/ 5X SSC (pH 4.5) at 65°C; Tris-HCl buffered saline /0.1% Tween(TBST)/2 mM levimasole at room temperature. Sections were then blocked in 5% sheep serum/ TBST, and incubated in anti-digoxigenin-AP Fab fragment antibody (Boehringer) at 4°C overnight. Sections were washed four times each in TBST, then in NTMT [100 mM NaCl, 100 mM Tris-HCl (pH 9.5), 50 mM MgCl_2 , 0.1% Tween, 2 mM levimasole and developed in 4-nitro blue tetrazolium chloride/5-bromo-4-chloro-3-indolyl-phosphate, dissolved in 10% polyvinyl alcohol/NTMT for 16 hr, and subsequently mounted.

Results

The *in situ* hybridization results demonstrate that NKX3.1 was expressed by several sexually dimorphic tissues of the male urogenital system, as well as by non-sexually dimorphic tissues including the dorsal aorta, lung, and intestine. Interestingly, NKX3.1 is primarily expressed by tissues that are derived from the endoderm, and is often found in epithelial cells that have a secretory function.

Within the male urogenital system, expression of NKX3.1 was observed in the developing urogenital sinus at day 14.5 *p.c.*, and in the ventral prostatic buds at day 17.5 *p.c.*. During the development of the male urogenital system, the primitive urogenital sinus is subdivided into three regions, which give rise to the urinary bladder, the prostatic and membranous parts of the urethra, and the penile urethra. Outbuddings of the central (pelvic) region of the urogenital sinus are responsible for formation of the prostate gland at late stages of embryogenesis. Thus, at day 14.5 of gestation, NKX3.1 expression was observed in outbuddings of the pelvic region of the urogenital sinus, with lower levels of expression in the prospective urethra. Notably, expression is confined to the epithelial cells that are invaginating into the surrounding mesenchyme, with highest levels observed at the leading edge. Later, at day 17.5 *p.c.*, expression of NKX3.1 is found in the developing ventral prostatic buds. NKX3.1 expression was also observed in the dorsolateral and anterior prostatic buds, but not in the epithelial lining of the bladder. In addition, NKX3.1 expression was also observed in the developing tests at days 14.5 and 17.5 of gestation. Expression was localized to the medullary cords, which form the seminiferous tubules, and was not observed in the interstitial mesenchyme or in the cells forming the fibrous outer layer of the testis. No expression of NKX3.1 was observed in the ductus deferens or in the epididymis.

Furthermore, NKX3.1 expression was observed in several non-sexually dimorphic tissues, such as in the endothelial cells that line the dorsal aorta at day 14.5 of gestation. Interestingly, NKX3.1 is not expressed by endothelial cells in general, but instead its expression is confined to regions of the dorsal aorta that

are in proximity to the developing metanephric kidneys, as well as to the renal arteries that arise from the dorsal aorta. In other non-sexually dimorphic tissues, low-level expression of NKX3.1 was observed in the epithelial cells that line the bronchi of the lung. Finally, expression of NKX3.1 was also observed in the small intestine, where expression is localized to a limited population of cells that appear to correspond to the mucus-secreting goblet cells.

Example 10: Temporal and Spatial Pattern of NKX3.1 Expression During Embryogenesis

To gain further insight into its potential developmental functions, the expression of NKX3.1 in pregastrulation through organogenesis stage embryos (e6.5-e10.5) and at a later stage of organogenesis (e15.5) was examined. These studies have revealed an intriguing pattern of expression that is markedly different from the Hox and other known homeobox genes, and implicate NKX3.1 in the differentiation of distinct populations of epithelial cells.

Materials and Methods

A 530-base ³⁵S-labeled antisense RNA probe derived from the 3' noncoding region of murine NKX3.1 was hybridized to serial paraffin sections of paraformaldehyde-fixed FVB/N embryos as described (Bogard *et al.*, 1989). The corresponding sense probe used on alternating sections did not hybridize in a specific pattern.

Results

To determine the temporal and spatial pattern of expression of NKX3.1 during embryogenesis, serial sections of mouse embryos at various stages were analyzed by *in situ* hybridization using an antisense RNA probe derived from the 3' non-coding region of the mRNA. Analysis of sections of e6.5 and e7.5 embryos did not reveal any hybridization above background, indicating that this gene is not activated during gastrulation. Prominent expression of NKX3.1 was first detected

in e8.5 embryos, and was localized to anterior paraxial mesenchyme. Mesenchyme adjacent to the developing midbrain and hindbrain expressed NKX3.1, with an apparent anterior boundary just posterior to the level of the foregut diverticulum. This head paraxial mesenchyme is the source of all
5 voluntary muscles of the head and, in addition, makes contributions to skeletal elements, the dermis, and meninges in the chick (Noden, D.M., *Am. J. Anat.* 168:257-276 (1983); Noden, D.M., *Develop. Biol.* 96:144-165 (1983)). Compared to somites which are patterned in large measure by Hox genes, little is known about the genetic mechanisms that specify the fate of anterior paraxial
10 mesenchyme. Expression of NKX3.1 in this tissue provides the first insight into potential pathways to determine cell specification in this tissue.

Expression of NKX3.1 was also seen in more mature anterior somites in e8.5 embryos, while newly-condensed somites and the unsegmented presomitic mesoderm did not show hybridization signal. This pattern of expression suggests
15 that NKX3.1 is activated in a cranio-caudal sequence, in a manner that parallels somite differentiation. Within the labeled somites, expression was not uniform, but was restricted to the ventral region. The onset of NKX3.1 expression in somites correlated well with the compartmentalization of the epithelial somite into the dermamyotome, which initially retains an epithelial arrangement, and the
20 sclerotome, which is clearly distinguishable by the mesenchymal phenotype of its cells, with NKX3.1 being confined to sclerotomal cells. This spatiotemporal pattern of expression is distinct from that of the Hox genes which are activated first in presomitic mesoderm, with expression spreading anteriorly to a discrete somitic boundary (Krumlauf, R., *Cell* 78:191-201 (1994)). The contrasting
25 patterns of Hox versus NKX3.1 expression in somites likely reflect different functions for these genes in the differentiation of paraxial mesoderm. The regionally-restricted, overlapping patterns of Hox gene expression may form the basis of a molecular "code" that is translated into specific vertebral phenotypes (Kessel and Gruss, 1991). Patterning functions attributed to Hox genes, such as
30 growth of ribs, have been suggested based on transplant studies to be established

prior to the onset of overt segmentation into somites (Kieny *et al.*, *Develop. Biol.* 28:142-161 (1972)). On the other hand, NKX3.1 may have a more general function in somite differentiation, for example by conferring a mesenchymal phenotype on somitic cells. The division of the somite into sclerotome and dermamyotome is an essential step in vertebral development that may be regulated by NKX3.1.

The pattern of expression at e9.5 was similar to that observed in e8.5 embryos. NKX3.1 mRNA continued to be expressed in the unsegmented paraxial mesenchyme anterior to the region of somite formation, but at a reduced level compared to e8.5, and in the sclerotome of differentiating somites. An anteroposterior gradient of expression was apparent in somites, with the more mature anterior somites showing less signal than those more posterior and just beginning to undergo differentiation. Highest levels of NKX3.1 expression appeared to correlate with early stages of sclerotome development, and expression was clearly absent from the dermamyotome. The down-regulation of NKX3.1 in more mature somites also argues for an early function in somite differentiation.

Analysis of e10.5 embryos revealed continued NKX3.1 expression in sclerotomal cells and head mesenchyme, as well as several new sites, most notably the axial mesenchyme. The axial mesenchyme consists of a loose meshwork of cells surrounding the notochord, directly beneath the neural tube, and is histologically distinct from the more dense paraxial mesenchyme from which somites derive (Verbout, A.J., "Advances in anatomy, embryology and cell biology," in *The Development of the Vertebral Column*, Vol. 90, Springer-Verlag, New York (1985)). NKX3.1 expression was observed in axial mesenchyme along the entire anteroposterior axis. Although these cells are contiguous with the sclerotomal cells, they are morphologically distinct, prompting the suggestion of different embryonic origins (Dawes, B., *Mus Musculus. philos. Trans. R. Soc. Lond. [Biol.]* 218:115-170 (1931)). More recent authors have argued for a common somitic origin for sclerotomal and axial mesenchyme, based solely on histologic criteria (Verbout, A.J., "Advances in anatomy, embryology and cell

biology," in *The Development of the Vertebral Column*, Vol. 90, Springer-Verlag, New York (1985)). Detection of NKX3.1 mRNA in both cell populations strongly supports the latter hypothesis by providing a genetic link between the two. NKX3.1 may play a role in specifying an axial mesenchyme fate in differentiating somitic cells.

A second new site of expression seen in e10.5 embryos is the wall of the abdominal aorta. The abdominal aorta develops in close apposition to the axial mesenchyme. In fact, there is no clear morphologic boundary which separates sclerotome, axial mesenchyme, and the wall of the aorta (Verbout, A.J., "Advances in anatomy, embryology and cell biology," in *The Development of the Vertebral Column*, Vol. 90, Springer-Verlag, New York (1985)). Scattered cells along the length of the abdominal aorta expressed NKX3.1 with an apparent anterior boundary just caudal to the level of the heart. These cells could represent axial mesenchymal cells that have been recruited to participate in vessel wall formation. Although the embryonic origin of endothelial cells has been well-characterized, the origin of vascular smooth muscle cells in developing vessels is poorly-defined and no markers that identify precursor cells have been reported (Owens, G.K., *Physiol. Rev.* 75:487-509 (1995)). NKX3.1 may represent an early marker of vascular smooth muscle cell differentiation in the abdominal aorta.

By e15.5, expression of NKX3.1 in somitic derivatives was no longer detectable. However, strong expression was observed in the abdominal aorta and in the developing renal vasculature. Although the limited cellular resolution of *in situ* hybridization precludes an unequivocal interpretation, NKX3.1 appears to be absent from the endothelial lining of the aorta. Immunostaining to detect smooth muscle α -actin, an early marker of smooth muscle cell differentiation (Owens, G.K., *Physiol. Rev.* 75:487-509 (1995)), revealed a perfect overlap with expression of NKX3.1 in these vessels in serial sections. Based on these observations, it appears that NKX3.1 is involved in specifying a smooth muscle cell phenotype in cells recruited from the axial mesenchyme into the wall of the aorta. Expression in the wall of the aorta was stronger posterior and faded out

towards the heart. Consistent with our observations on e10.5 embryos, NKX3.1 was not detected in vessels anterior to the level of the heart.

Prominent expression of NKX3.1 was also detected in the developing incisors and molars in e15.5 embryos. The first evidence of tooth development is a thickening of the oral epithelium detectable in e12.5 embryos (Cohn, S.A., *Am. J. Anat.* 101:295-320 (1957)). This dental lamina develops into an epithelial bud that becomes surrounded by mesenchymal cells derived from the neural crest that constitute the dental papilla. By e15.5, invaginations of the dental lamina mark the beginning of the cap stage of development. The dental lamina also begins to differentiate into two distinct layers, the inner and outer enamel layers. NKX3.1 is expressed predominantly in the deep invaginations of the dental lamina at the junction between the inner and outer enamel layers. The surrounding mesenchyme of the dental papilla did not express NKX3.1. This restricted pattern of expression in the developing teeth suggests that NKX3.1 is involved in regional differentiation of cells within the dental epithelium.

The presence of a homeobox in the coding region of NKX3.1 provides presumptive evidence that this gene encodes a transcription factor that, by analogy to other homeodomain proteins, likely regulates genes involved in establishing a specific state of cellular differentiation. The onset of expression of NKX3.1 in somites seems to occur in concert with the transition from an entirely epithelial structure to one with two morphologically distinct components; it is possible that NKX3.1 may regulate the epithelial to mesenchymal transition that gives rise to the sclerotomal cell population where it is expressed. The observation of NKX3.1 expression in axial mesoderm and the wall of the abdominal aorta may indicate that NKX3.1 is involved in directing a subset of somitic cells toward a particular cell fate. The distribution of NKX3.1 in specific regions of developing teeth is also consistent with a role in the commitment of epithelial cells.

It will be clear that the invention may be practiced otherwise than as particularly described in the foregoing description and examples.

Numerous modifications and variations of the present invention are possible in light of the above teachings and, therefore, are within the scope of the appended claims.

5 The entire disclosure of all publications (including patents, patent applications, journal articles, laboratory manuals, books, or other documents) cited herein are hereby incorporated by reference.

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SEQUENCE LISTING

(1) GENERAL INFORMATION:

(i) APPLICANT: HUMAN GENOME SCIENCES, INC.
9410 KEY WEST AVENUE
ROCKVILLE, MD 20850
UNITED STATES OF AMERICA

APPLICANT/INVENTOR: Carter, Kenneth C.
He, Wei-Wu

(ii) TITLE OF INVENTION: Human NK-3 Related Prostate Specific
Gene-1

(iii) NUMBER OF SEQUENCES: 30

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(D) STATE: DC
(E) COUNTRY: USA
(F) ZIP: 20005

(v) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: Floppy disk
(B) COMPUTER: IBM PC compatible
(C) OPERATING SYSTEM: PC-DOS/MS-DOS
(D) SOFTWARE: PatentIn Release #1.0, Version #1.30

(vi) CURRENT APPLICATION DATA:

(A) APPLICATION NUMBER: To be assigned
(B) FILING DATE: Herewith
(C) CLASSIFICATION:

(vii) PRIOR APPLICATION DATA:

(A) APPLICATION NUMBER: US 60/051,080
(B) FILING DATE: 27-JUN-1997

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(B) REGISTRATION NUMBER: 36,688
(C) REFERENCE/DOCKET NUMBER: 1488.079PC01

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(B) TELEFAX: 301-309-8439

(2) INFORMATION FOR SEQ ID NO:1:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 705 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

-89-

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

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CGCGCCGCGC CGGAGGAGGC CGAGACGCTG GCAGAGACCG AGCCAGAAAG GCACTTGGGG      300
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AAGCAGCCGC AGAAGCGCTC CCGAGCTGCC TTCTCCACA CTCAGGTGAT CGAGTTGGAG      420
AGGAAGTTCA GCCATCAGAA GTACCTGTCG GCCCCTGAAC GGGCCACCT GGCCAAGAAC      480
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CGAAAGCAGC TCTCTCGGA GCTGGGAGAC TTGGAGAAGC ACTCCTCTTT GCCGGCCCTG      600
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(2) INFORMATION FOR SEQ ID NO:2:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 234 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

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Asp Ile Leu Arg Asp Gly Ala Gln Arg Gln Gly Gly Arg Thr Ser Ser
35           40           45
Gln Arg Gln Arg Asp Pro Glu Pro Glu Pro Glu Pro Glu Gly
50           55           60
Gly Arg Ser Arg Ala Gly Ala Gln Asn Asp Gln Leu Ser Thr Gly Pro
65           70           75           80
Arg Ala Ala Pro Glu Glu Ala Glu Thr Leu Ala Glu Thr Glu Pro Glu
85           90           95
Arg His Leu Gly Ser Tyr Leu Leu Asp Ser Glu Asn Thr Ser Gly Ala
100          105          110
Leu Pro Arg Leu Pro Gln Thr Pro Lys Gln Pro Gln Lys Arg Ser Arg

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-90-

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145	150	155
Leu Lys Leu Thr Glu Thr Gln Val Lys Ile Trp Phe Gln Asn Arg Arg		
	165	170
Tyr Lys Thr Lys Arg Lys Gln Leu Ser Ser Glu Leu Gly Asp Leu Glu		
	180	185
Lys His Ser Ser Leu Pro Ala Leu Lys Glu Glu Ala Phe Ser Arg Ala		
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Cys Val Gly Ser Trp Ser Pro Ala Phe Gly		
225	230	

(2) INFORMATION FOR SEQ ID NO:3:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 705 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

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CGAAAGCAGC TCTCCTCGGA GCTGGGAGAC TTGGAGAAGC ACTCCTCTTT GCCGGCCCTG	600
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(2) INFORMATION FOR SEQ ID NO:4:

-91-

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 234 amino acids
 (B) TYPE: amino acid
 (C) STRANDEDNESS: single
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

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35           40           45
Gln Arg Gln Cys Asp Pro Glu Pro Glu Pro Glu Pro Glu Pro Glu Gly
50           55           60
Gly Arg Ser Arg Ala Gly Ala Gln Asn Asp Gln Leu Ser Thr Gly Pro
65           70           75           80
Arg Ala Ala Pro Glu Glu Ala Glu Thr Leu Ala Glu Thr Glu Pro Glu
85           90           95
Arg His Leu Gly Ser Tyr Leu Leu Asp Ser Glu Asn Thr Ser Gly Ala
100          105          110
Leu Pro Arg Leu Pro Gln Thr Pro Lys Gln Pro Gln Lys Arg Ser Arg
115          120          125
Ala Ala Phe Ser His Thr Gln Val Ile Glu Leu Glu Arg Lys Phe Ser
130          135          140
His Gln Lys Tyr Leu Ser Ala Pro Glu Arg Ala His Leu Ala Lys Asn
145          150          155          160
Leu Lys Leu Thr Glu Thr Gln Val Lys Ile Trp Phe Gln Asn Arg Arg
165          170          175
Tyr Lys Thr Lys Arg Lys Gln Leu Ser Ser Glu Leu Gly Asp Leu Glu
180          185          190
Lys His Ser Ser Leu Pro Ala Leu Lys Glu Glu Ala Phe Ser Arg Ala
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210          215          220
Cys Val Gly Ser Trp Ser Pro Ala Phe Gly
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(2) INFORMATION FOR SEQ ID NO:5:

- (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 60 amino acids
 (B) TYPE: amino acid

-92-

(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

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20 25 30
His Leu Ala Lys Asn Leu Lys Leu Thr Glu Thr Gln Val Lys Ile Trp
35 40 45
Phe Gln Asn Arg Arg Tyr Lys Thr Lys Arg Lys Gln
50 55 60

(2) INFORMATION FOR SEO ID NO: 6:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 60 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

Lys	Lys	Arg	Ser	Arg	Ala	Ala	Phe	Ser	His	Ala	Gln	Val	Phe	Glu	Leu
1				5					10					15	
Glu	Arg	Arg	Phe	Ala	Gln	Gln	Arg	Tyr	Leu	Ser	Gly	Pro	Glu	Arg	Ser
			20					25					30		
Glu	Met	Ala	Lys	Ser	Leu	Arg	Leu	Thr	Glu	Thr	Gln	Val	Lys	Ile	Trp
		35					40					45			
Phe	Gln	Asn	Arg	Arg	Tyr	Lys	Thr	Lys	Arg	Lys	Gln				
	50					55					60				

(2) INFORMATION FOR SEQ ID NO:7:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 60 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

Lys Lys Arg Lys Arg Arg Val Leu Phe Thr Lys Ala Gln Thr Tyr Glu

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1             5             10             15
Leu Glu Arg Arg Phe Arg Gln Gln Arg Tyr Leu Ser Ala Pro Glu Arg
                20                25                30
Glu His Leu Ala Ser Leu Ile Arg Leu Thr Pro Thr Gln Val Lys Ile
                35                40                45
Trp Phe Gln Asn His Arg Tyr Lys Thr Lys Arg Ala
                50                55                60

```

(2) INFORMATION FOR SEQ ID NO:8:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 60 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

```

Lys Arg Lys Pro Arg Val Leu Phe Ser Gln Ala Gln Val Leu Glu Leu
1             5             10             15
Glu Cys Arg Phe Arg Leu Lys Lys Tyr Leu Thr Gly Ala Glu Arg Glu
                20                25                30
Ile Ile Ala Gln Lys Leu Asn Leu Ser Ala Thr Gln Val Lys Ile Trp
                35                40                45
Phe Gln Asn Arg Arg Tyr Lys Ser Lys Arg Gly Asp
                50                55                60

```

(2) INFORMATION FOR SEQ ID NO:9:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 124 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

```

Met Leu Arg Val Ala Glu Pro Arg Glu Pro Arg Val Glu Ala Gly Gly
1             5             10             15
Arg Ser Pro Trp Ala Ala Pro Pro Thr Gln Ser Lys Arg Leu Thr Ser
                20                25                30
Phe Leu Ile Gln Asp Ile Leu Arg Asp Arg Ala Glu Arg His Gly Gly
                35                40                45
His Ser Gly Asn Pro Gln His Ser Pro Asp Pro Arg Arg Asp Ser Ala

```

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50		55		60											
Pro	Glu	Pro	Asp	Lys	Ala	Gly	Gly	Arg	Gly	Val	Ala	Pro	Glu	Asp	Pro
65					70					75					80
Pro	Ser	Ile	Arg	His	Ser	Pro	Ala	Glu	Thr	Pro	Thr	Glu	Pro	Glu	Ser
			85						90					95	
Asp	Ala	His	Phe	Glu	Thr	Tyr	Leu	Leu	Asp	Cys	Glu	His	Asn	Pro	Gly
		100						105					110		
Asp	Leu	Ala	Ser	Ala	Pro	Gln	Val	Thr	Lys	Gln	Pro				
		115						120							

(2) INFORMATION FOR SEQ ID NO:10:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 122 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

Met	Leu	Arg	Val	Pro	Glu	Pro	Arg	Pro	Gly	Glu	Ala	Lys	Ala	Glu	Gly
1				5					10					15	
Ala	Ala	Pro	Pro	Thr	Pro	Ser	Lys	Pro	Leu	Thr	Ser	Phe	Leu	Ile	Asp
			20					25					30		
Ile	Leu	Arg	Asp	Gly	Ala	Gln	Arg	Gln	Gly	Gly	Arg	Thr	Ser	Ser	Gln
		35					40					45			
Arg	Gln	Cys	Asp	Pro	Glu	Pro	Glu	Pro	Glu	Pro	Glu	Pro	Glu	Gly	Gly
	50					55					60				
Arg	Ser	Arg	Ala	Gly	Ala	Gln	Asn	Asp	Gln	Leu	Ser	Thr	Gly	Pro	Arg
65				70						75					80
Ala	Ala	Pro	Glu	Glu	Ala	Glu	Thr	Leu	Ala	Glu	Thr	Glu	Pro	Glu	Arg
			85					90						95	
His	Leu	Gly	Ser	Tyr	Leu	Leu	Asp	Ser	Glu	Asn	Thr	Ser	Gly	Ala	Leu
		100						105					110		
Pro	Arg	Leu	Pro	Gln	Thr	Pro	Lys	Gln	Pro						
		115						120							

(2) INFORMATION FOR SEQ ID NO:11:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 60 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

Gln	Lys	Arg	Ser	Arg	Ala	Ala	Phe	Ser	His	Thr	Gln	Val	Ile	Glu	Leu
1				5					10					15	
Glu	Arg	Lys	Phe	Ser	His	Gln	Lys	Tyr	Leu	Ser	Ala	Pro	Glu	Arg	Ala
			20					25					30		
His	Leu	Ala	Lys	Asn	Leu	Lys	Leu	Thr	Glu	Thr	Gln	Val	Lys	Ile	Trp
		35					40					45			
Phe	Gln	Asn	Arg	Arg	Tyr	Lys	Thr	Lys	Arg	Lys	Gln				
	50					55					60				

(2) INFORMATION FOR SEQ ID NO:12:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 60 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

Gln	Lys	Arg	Ser	Arg	Ala	Ala	Phe	Ser	His	Thr	Gln	Val	Ile	Glu	Leu
1				5					10					15	
Glu	Arg	Lys	Phe	Ser	His	Gln	Lys	Tyr	Leu	Ser	Ala	Pro	Glu	Arg	Ala
			20					25					30		
His	Leu	Ala	Lys	Asn	Leu	Lys	Leu	Thr	Glu	Thr	Gln	Val	Lys	Ile	Trp
		35					40					45			
Phe	Gln	Asn	Arg	Arg	Tyr	Lys	Thr	Lys	Arg	Lys	Gln				
	50					55					60				

(2) INFORMATION FOR SEQ ID NO:13:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 53 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

Leu	Ser	Glu	Asp	Leu	Gly	Val	Leu	Glu	Lys	Asn	Ser	Pro	Leu	Ser	Leu
1				5					10					15	
Pro	Ala	Leu	Lys	Asp	Asp	Ser	Leu	Pro	Ser	Thr	Ser	Leu	Val	Ser	Val
			20					25					30		

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Tyr Thr Ser Tyr Pro Tyr Tyr Pro Tyr Leu Tyr Cys Leu Gly Ser Trp
 35 40 45

His Pro Ser Phe Trp
 50

(2) INFORMATION FOR SEQ ID NO:14:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 51 amino acids
 - (B) TYPE: amino acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:

Leu Ser Ser Glu Leu Gly Asp Leu Glu Lys His Ser Ser Leu Pro Ala
 1 5 10 15

Leu Lys Glu Glu Ala Phe Ser Arg Ala Ser Leu Val Ser Val Tyr Asn
 20 25 30

Ser Tyr Pro Tyr Tyr Pro Tyr Leu Tyr Cys Val Gly Ser Trp Ser Pro
 35 40 45

Ala Phe Gly
 50

(2) INFORMATION FOR SEQ ID NO:15:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 27 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:

GCGGGATCCA TGCTCAGGGT TCCGGAG

27

(2) INFORMATION FOR SEQ ID NO:16:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 26 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:

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GCGAGCTTTT ACCCAAAAGC TGGGCT

26

(2) INFORMATION FOR SEQ ID NO:17:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 28 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

GCGGGATCCC ATGCTCAGGG TTCCGAG

28

(2) INFORMATION FOR SEQ ID NO:18:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 26 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:

GCGGATCCTT ACCCAAAAGC TGGGCT

26

(2) INFORMATION FOR SEQ ID NO:19:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 26 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:

GCGGATCCTT ACCCAAAAGC TGGGCT

26

(2) INFORMATION FOR SEQ ID NO:20:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 24 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:

CACTGCCCAG TCAAGTGTTT TTGA

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(2) INFORMATION FOR SEQ ID NO:21:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 24 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:

CACTGCCCAG TCACGTGTTT GTGA

24

(2) INFORMATION FOR SEQ ID NO:22:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 15 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:

ACACTAATTG GAGGC

15

(2) INFORMATION FOR SEQ ID NO:23:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 15 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:

ACACTACTTG GAGGC

15

(2) INFORMATION FOR SEQ ID NO:24:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 19 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:24:

CTCTAATGGC TTTTTTCTC

19

(2) INFORMATION FOR SEQ ID NO:25:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 3974 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: both

(D) TOPOLOGY: both

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:25:

GGTACCTAAG TGAGTAGGGC GTCCGATCGA CGGACGCCTT TTTTTTGAAT TCGTAATCAT	60
GGTCATAGCT GTTTCCTGTG TGAAATTGTT ATCCGCTCAC AATTCCACAC AACATACGAG	120
CCGGAAGCAT AAAGTGTAAG GCCTGGGGTG CCTAATGAGT GAGCTAACTC ACATTAATTG	180
CGTTGCGCTC ACTGCCCCTG TTCCAGTCGG GAAACCTGTC GTGCCAGCTG CATTAAATGAA	240
TCGGCCAACG CGCGGGGAGA GGCGGTTTGC GTATTGGGCG CTCTTCCGCT TCCTCGCTCA	300
CTGACTCGCT GCGCTCGGTC GTTCGGCTGC GGCGAGCGGT ATCAGCTCAC TCAAAGGCGG	360
TAATACGGTT ATCCACAGAA TCAGGGGATA ACGCAGGAAA GAACATGTGA GCAAAGGCC	420
AGCAAAAGGC CAGGAACCGT AAAAAGGCCG CGTTGCTGGC GTTTTCCAT AGGCTCCGCC	480
CCCCTGACGA GCATCAGAAA AATCGACGCT CAAGTCAGAG GTGGCGAAAC CCGACAGGAC	540
TATAAAGATA CCAGGCGTTT CCCCTGGAA GCTCCCTCGT GCGCTCTCCT GTTCCGACCC	600
TGCCGCTTAC CGGATACCTG TCCGCCTTTC TCCCTTCGGG AAGCGTGGCG CTTTCTCATA	660
GCTCACGCTG TAGGTATCTC AGTTCGGTGT AGGTCGTTTC CTCCAAGCTG GGCTGTGTGC	720
ACGAACCCCC CGTTCAGCCC GACCGCTGCG CCTTATCCGG TAACTATCGT CTTGAGTCCA	780
ACCCGGTAAG ACACGACTTA TCGCCACTGG CAGCAGCCAC TGGTAACAGG ATTAGCAGAG	840
CGAGGTATGT AGGCGGTGCT ACAGAGTTCT TGAAGTGGTG GCCTAACTAC GGCTACACTA	900
GAAGAACAGT ATTTGGTATC TGCGCTCTGC TGAAGCCAGT TACCTTCGGA AAAAGAGTTG	960
GTAGCTCTTG ATCCGGCAAA CAAACCACCG CTGGTAGCGG TGGTTTTTTT GTTTGCAAGC	1020
AGCAGATTAC GCGCAGAAAA AAAGGATCTC AAGAAGATCC TTTGATCTTT TCTACGGGGT	1080
CTGACGCTCA GTGGAACGAA AACTCACGTT AAGGGATTTT GGTCATGAGA TTATCGTCGA	1140
CAATTCGCGC GCGAAGGCGA AGCGGCATGC ATTTACGTTG ACACCATCGA ATGGTGCAAA	1200
ACCTTTCGCG GTATGGCATG ATAGCGCCCG GAAGAGAGTC AATTCAGGGT GGTGAATGTG	1260

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AAACCAGTAA CGTTATACGA TGTCGCAGAG TATGCCGGTG TCTCTTATCA GACCGTTTCC	1320
CGCGTGGTGA ACCAGGCCAG CCACGTTTCT GCGAAAACGC GGGAAAAAGT GGAAGCGGCG	1380
ATGGCGGAGC TGAATTACAT TCCCAACCGC GTGGCACAAC AACTGGCGGG CAAACAGTCG	1440
TTGCTGATTG GCGTTGCCAC CTCCAGTCTG GCCCTGCACG CGCCGTCGCA AATTGTCGCG	1500
GCGATTAAAT CTCGCGCCGA TCAACTGGGT GCCAGCGTGG TGGTGTGAT GGTAGAACGA	1560
AGCGGCGTCG AAGCCTGTAA AGCGGCGGTG CACAATCTTC TCGCGCAACG CGTCAGTGGG	1620
CTGATCATTG ACTATCCGCT GGATGACCAG GATGCCATTG CTGTGGAAGC TGCCTGCACT	1680
AATGTTCCGG CGTTATTTCT TGATGTCTCT GACCAGACAC CCATCAACAG TATTATTTTC	1740
TCCCATGAAG ACGGTACGCG ACTGGGCGTG GAGCATCTGG TCGCATTGGG TCACCAGCAA	1800
ATCGCGCTGT TAGCGGGCCC ATTAAGTTCT GTCTCGGCGC GTCTGCGTCT GGCTGGCTGG	1860
CATAAATATC TCACTCGCAA TCAAATTCAG CCGATAGCGG AACGGGAAGG CGACTGGAGT	1920
GCCATGTCCG GTTTTCAACA AACCATGCAA ATGCTGAATG AGGGCATCGT TCCCACTGCG	1980
ATGCTGGTTG CCAACGATCA GATGGCGCTG GCGCAATGC GCGCCATTAC CGAGTCCGGG	2040
CTGCGCGTTG GTGCGGATAT CTCGGTAGTG GGATACGACG ATACCGAAGA CAGCTCATGT	2100
TATATCCCGC CGTTAACCAC CATCAAACAG GATTTTCGCC TGCTGGGGCA AACCAGCGTG	2160
GACCGCTTGC TGCAACTCTC TCAGGGCCAG GCGGTGAAGG GCAATCAGCT GTTGCCCGTC	2220
TCACTGGTGA AAAGAAAAAC CACCCTGGCG CCCAATACGC AAACCGCCTC TCCCCGCGCG	2280
TTGGCCGATT CATTAAATGCA GCTGGCACGA CAGGTTTCCC GACTGGAAG CGGGCAGTGA	2340
GCGCAACGCA ATTAATGTAA GTTAGCGCGA ATTGTGACG AAAGCGGCCA TCGTGCCTCC	2400
CCACTCCTGC AGTTCGGGGG CATGGATGCG CGGATAGCCG CTGCTGGTTT CCTGGATGCC	2460
GACGGATTTG CACTGCCGGT AGAACTCCGC GAGGTCGTCC AGCCTCAGGC AGCAGCTGAA	2520
CCAACTCGCG AGGGGATCGA GCGCGGGTG GCGGAAGAAC TCCAGCATGA GATCCCCGCG	2580
CTGGAGGATC ATCCAGCCGG CGTCCCGGAA AACGATTCCG AAGCCCAACC TTTCATAGAA	2640
GGCGGCGGTG GAATCGAAAT CTCGTGATGG CAGGTTGGGC GTCGCTTGGT CGGTCATTTT	2700
GAACCCAGAG GTCCCGCTCA GAAGAACTCG TCAAGAAGGC GATAGAAGGC GATGCGCTGC	2760
GAATCGGGAG CGGCGATACC GTAAAGCACG AGGAAGCGGT CAGCCCATTG GCCGCCAAGC	2820
TCTTCAGCAA TATCACGGGT AGCCAACGCT ATGTCCTGAT AGCGGTCCGC CACACCCAGC	2880
CGGCCACAGT CGATGAATCC AGAAAAGCGG CCATTTTCCA CCATGATATT CGGCAAGCAG	2940
GCATCGCCAT GGGTCACGAC GAGATCCTCG CCGTCGGGCA TGCGCGCCTT GAGCCTGGCG	3000
AACAGTTCGG CTGGCGCGAG CCCCTGATGC TCTTCGTCCA GATCATCCTG ATCGACAAGA	3060
CCGGCTTCCA TCCGAGTACG TGCTCGCTCG ATGCGATGTT TCGCTTGGTG GTCGAATGGG	3120

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CAGGTAGCCG GATCAAGCGT ATGCAGCCGC CGCATTGCAT CAGCCATGAT GGATACTTTC	3180
TCGGCAGGAG CAAGGTGAGA TGACAGGAGA TCCTGCCCCG GCACTTCGCC CAATAGCAGC	3240
CAGTCCCTTC CCGCTTCAGT GACAACGTCG AGCACAGCTG CGCAAGGAAC GCCCGTCGTG	3300
GCCAGCCACG ATAGCCGCGC TGCCTCGTCC TGCAGTTCAT TCAGGGCACC GGACAGGTCG	3360
GTCTTGACAA AAAGAACCGG GCGCCCCTGC GCTGACAGCC GGAACACGGC GGCATCAGAG	3420
CAGCCGATTG TCTGTTGTGC CCAATCATAG CCGAATAGCC TCTCCACCCA AGCGGCCGGA	3480
GAACCTGCGT GCAATCCATC TTGTTCAATC ATGCGAAACG ATCCTCATCC TGTCTCTTGA	3540
TCAGATCTTG ATCCCCTGCG CCATCAGATC CTTGGCGGCA AGAAAGCCAT CCAGTTTACT	3600
TTGCAGGGCT TCCCAACCTT ACCAGAGGGC GCCCCAGCTG GCAATTCCGG TTCGCTTGCT	3660
GTCCATAAAA CCGCCAGTC TAGCTATCGC CATGTAAGCC CACTGCAAGC TACCTGCTTT	3720
CTCTTTGCGC TTGCGTTTTT CCTTGTCCAG ATAGCCAGT AGCTGACATT CATCCGGGGT	3780
CAGCACCGTT TCTGCGGACT GGCTTTCTAC GTGTTCCGCT TCCTTTAGCA GCCCTTGCGC	3840
CCTGAGTGCT TGCGGCAGCG TGAAGCTTAA AAACTGCAA AAAATAGTTT GACTTGTGAG	3900
CGGATAACAA TTAAGATGTA CCCAATTGTG AGCGGATAAC AATTTCACAC ATTAAAGAGG	3960
AGAAATTACA TATG	3974

(2) INFORMATION FOR SEQ ID NO:26:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 112 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: both
 - (D) TOPOLOGY: both

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:26:

AAGCTTAAAA AACTGCAAAA AATAGTTTGA CTTGTGAGCG GATAACAATT AAGATGTACC	60
CAATTGTGAG CGGATAACAA TTTCACACAT TAAAGAGGAG AAATTACATA TG	112

(2) INFORMATION FOR SEQ ID NO:27:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 415 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: both
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:27:

GCTCGAGCTT TCCACGCAAA TCTCCAATAG CGTTGGTGTG CACACTCTTG CTCCCTGACC	60
TTGGCCCCCTT GTGCTTTTCA TAGAGACACC CTGGGGAAGG CAGTTTTTGA TCATAATGGT	120
TGTCACCTGA GCTGGCATT A CAAAAAGCT GGGCTCCACT GCCACGCAG TACAGGTATG	180
GGTAGTAAGG ATAGCTGTTA TACACGGAGA CCAGGGAGGC CCGGAGAAG GCCTCTCTTT	240
CAGGGCCGGC AAAGAGGAGT GCTTCTCCAA GTCTCCAGC TCCGAGGAGA GCTGCTTTCG	300
CTTAGTCTTA TAGCGTCTGT TCTGGAACCA TATCTTCACT TGGGTCTCCG TGAGCTTGAG	360
GTTCTTGCC AGGTGGGCG TTCAGGGCGA TCAGGTACTT CTGATGGCTG AACTT	415

(2) INFORMATION FOR SEQ ID NO:28:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 507 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: both
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:28:

TTTTTTTCCC GTCCGACCCT CCGGAGTTTA TCACTTCCA GCGGTACTGG GCTGGACCGA	60
GCCTTGGTGG CATCCGATTC AGCTCCAGCA GCATCCCAGG TCCTATCCAG CATGGGCGGG	120
GCACCGCTGG TCAGAGCTCT GGCCTCAGG AATCTTCGGA CTCGTCCCTT TCCTCGAAGT	180
CGGGCTCGGG CTCTGGGTCC GGTCTGGCT CCAGTCCGG CTCTGGTTCT GCTTCAGCCT	240
CGGGTTCCAG CTCTGGCTCT GGCTCCGGCT CCGGTCAGC CTCTTAGAG GCCTCAGCTT	300
CCAGCTCTGG CTCTGGCTCC TCCGGGTCC CGGATTGCAC CGCTCGGAG CGCTCAGGCC	360
CTCCGGCTAC GTCAGGTCT GAAGCTCTGG GCAACCCGGG GCATGTGGTC ACCGAGCCGG	420
GTGCCTCCGG GTCCAGTGGC CTGGGCATGG AGGGTCATAG CTGCGGTCGC GGTAGCCCGG	480
ACCCACGTAC TGCGAGTCGA AAGCGGG	507

(2) INFORMATION FOR SEQ ID NO:29:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 533 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: both
 - (D) TOPOLOGY: both

(ii) MOLECULE TYPE: cDNA

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:29:

NGCACAGCAG NNAGGCACTT GGAGGTCTTA TCTGTTNGAC TTGNTNAACA CTTCAGGCGC	60
CCTTCCAAGG CTTCCCCAAA CCCCTAAGCA GCCGCANAAG CGCTCCCGAG CTGCCTTCTC	120
CCCACTCAG GTGATCGAGT TGGAGAGGAA GTTCAGCCAT CAGAAGTACC TGTCGGCCCC	180
TGGAACGGGC CCACCTGGCC AAGAACCTCA AGCTCACGGA GACCCAAGTG AAGATATGGT	240
TCCAGAACAG ACGCTATAAG ACTAAGCGAA ACAGCTCTCC TCGGANCTGG GGAGACTTGG	300
AGAAGCACTC CTCTTTGNCC GGCCCTGAAN GAGGAGGCCN TCNCCCGGGC CNNCCTGGTC	360
NCCGTGTAAT AACAGCGANC CTTNANTACC CATACTGTGA ANGCTGGGG CANNGGGAAG	420
CCCAGTTTNT TGGTAAANGG CCAGCTCCAG GTGAACAACC ATTAAGGATC AAAANGGCCT	480
TNCCCCAGGG TGTTNCATGG AAAAGCACAA GGGGCAAGGT CAGGAGCAA AGN	533

(2) INFORMATION FOR SEQ ID NO:30:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 318 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: both
 - (D) TOPOLOGY: both

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:30:

GGGAGCTGGA GCCAGAGCCA GAGCCAGATC TCAGGAAACT GGGGTCATTG CATAGAGGCT	60
GCCAGACAGT CTGCAGAGCT CAGCGGCCTG GGTTCAAACC TTCTCGCACA CTGCCACTGT	120
CGGTTACTTT GGCTTTCTAG AGCCAGATTC CTTGGCCATG AAATGGGTAC TGCTTACTTC	180
CCAGGTTATT TTGAGAATGA AGTGAGATGA AGTCAACAGT AGATGTATCT GTCCGTTGTC	240
CCTGCCCTGC TGTGGGGATG ACAGAGTGAT TTTGGACAAG ACCCAAGGNC TCGCTGGGCA	300
TCACTGGTCT TTCTTCAG	318

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Applicant's or agent's file reference number 1488.079PC01	International application No. To be assigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM
(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>7</u> , line <u>8</u> .	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	
formerly at 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit 28 April 1997	Accession Number 209005
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
DNA Plasmid NKX3.1	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable) The indications listed below will be submitted to the international Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

For receiving Office use only	For International Bureau use only
<input checked="" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer <i>Melvin Brooks</i>	Authorized officer

-103/2-

Applicant's or agent's file reference number 1488.079PC01	International application No. To be assigned
--	---

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM
(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>7</u> , line <u>13</u> .	
B. IDENTIFICATION OF DEPOSIT	
Further deposits are identified on an additional sheet <input checked="" type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country)	
10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	formerly at 12301 Parklawn Drive Rockville, Maryland 20852 United States of America
Date of deposit 28 April 1997	Accession Number 209006
C. ADDITIONAL INDICATIONS (leave blank if not applicable)	
This information is continued on an additional sheet <input type="checkbox"/>	
DNA Plasmid HPFCA 19	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the international Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

For receiving Office use only	For International Bureau use only
<input checked="" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer <i>Melvin Brooks</i>	Authorized officer

-103/3-

Applicant's or agent's file reference number 1488.079PC01	International application No. 1488.079PC01 assigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM
(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>19</u> , line <u>6</u> .	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 10801 University Boulevard formerly at 12301 Parklawn Drive Manassas, Virginia 20110-2209 Rockville, Maryland 20852 United States of America United States of America	
Date of deposit 25 February 1998	Accession Number 209645
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
DNA Plasmid pHE4a	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the international Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	

For receiving Office use only	For International Bureau use only
<input checked="" type="checkbox"/> This sheet was received with the international application	<input type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer <i>Melvin Brooks</i>	Authorized officer

What Is Claimed Is:

1. An isolated nucleic acid molecule selected from the group consisting of:

1. a polynucleotide encoding a polypeptide comprising amino acids from about 1 to about 234 in SEQ ID NOs:2 or 4;

- (b) a polynucleotide encoding a polypeptide comprising amino acids from about 2 to about 234 in SEQ ID NOs:2 or 4;

- (c) a polynucleotide encoding a polypeptide comprising amino acids from about 123 to about 153 in SEQ ID NOs:2 or 4;

- (d) a polynucleotide encoding a polypeptide comprising the amino acid sequence encoded by the cDNA clone contained in ATCC Deposit No. 209005 or 209006;

- (e) the complement of (a), (b), (c), or (d);

- (f) a polynucleotide variant created by altering the polynucleotide of (a), wherein:

- (1) said altering includes a nucleotide insertion, deletion, or substitution, or any combination thereof; and

- (2) the number of alterations is equal to or less than 5 % of the total number of nucleotides present in (a);

- (h) a polynucleotide variant created by altering the polynucleotide of (b), wherein:

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- 5
- (1) said altering includes a nucleotide insertion, deletion, or substitution, or any combination thereof; and
- (2) the number of alterations is equal to or less than 5 % of the total number of nucleotides present in (b);
- (i) a polynucleotide variant created by altering the polynucleotide of (c), wherein:
- 10
- (1) said altering includes a nucleotide insertion, deletion, or substitution, or any combination thereof; and
- (2) the number of alterations is equal to or less than 5 % of the total number of nucleotides present in (c);
- 15
- (j) a polynucleotide variant created by altering the polynucleotide of (d), wherein:
- (1) said altering includes a nucleotide insertion, deletion, or substitution, or any combination thereof; and
- 20
- (2) the number of alterations is equal to or less than 5 % of the total number of nucleotides present in (d); and
- (k) a polynucleotide variant created by altering the polynucleotide of (e), wherein:
- 25
- (1) said altering includes a nucleotide insertion, deletion, or substitution, or any combination thereof; and
- (2) the number of alterations is equal to or less than 5 % of the total number of nucleotides present in (e).
- 30

2. The nucleic acid molecule of claim 1 wherein said polynucleotide has the complete nucleotide sequence in SEQ ID NOs:1, 3 or 8.

5 3. The nucleic acid molecule of claim 1 wherein said polynucleotide has the nucleotide sequence in SEQ ID NOs:1, 3 or 8 encoding the NKX3.1 polypeptide having the complete amino acid sequence in SEQ ID NOs:2 or 4.

4. The nucleic acid molecule of claim 1 wherein said polynucleotide has the complete nucleotide sequence of the cDNA clone contained in ATCC Deposit No 209005 or 209006.

10 5. The nucleic acid molecule of claim 1 wherein said polynucleotide has the nucleotide sequence encoding the NKX3.1 polypeptide having the complete amino acid sequence encoded by the cDNA clone contained in ATCC Deposit No. 209005 or 209006.

15 6. An isolated nucleic acid molecule comprising a polynucleotide which hybridizes under stringent hybridization conditions to a polynucleotide having a nucleotide sequence identical to a nucleotide sequence in (a), (b), (c), (d), (e), (f) or (g) of claim 1 wherein said polynucleotide which hybridizes does not hybridize under stringent hybridization conditions to a polynucleotide having a nucleotide sequence consisting of only A residues or of only T residues.

20 7. An isolated nucleic acid molecule comprising a polynucleotide which encodes the amino acid sequence of an epitope-bearing portion of a NKX3.1 polypeptide having an amino acid sequence in (a), (b), (c), (d), (e) or (f) of claim 1.

8. The isolated nucleic acid molecule of claim 9, which encodes an epitope-bearing portion of a NKX3.1 polypeptide selected from the group consisting of: a polypeptide comprising amino acid residues from about 1 to about 13 in SEQ ID NO:2 or SEQ ID NO:4; a polypeptide comprising amino acid residues from about 18 to about 24 in SEQ ID NO:2 or SEQ ID NO:4; a polypeptide comprising amino acid residues from about 35 to about 89 in SEQ ID NO:2 or SEQ ID NO:4; a polypeptide comprising amino acid residues from about 103 to about 113 in SEQ ID NO:2 or SEQ ID NO:4; a polypeptide comprising amino acid residues from about 117 to about 130 in SEQ ID NO:2 or SEQ ID NO:4; and a polypeptide comprising amino acid residues from about 173 to about 185 in SEQ ID NO:2 or SEQ ID NO:4.

9. A method for making a recombinant vector comprising inserting an isolated nucleic acid molecule of claim 1 into a vector.

10. A recombinant vector produced by the method of claim 9.

11. A method of making a recombinant host cell comprising introducing the recombinant vector of claim 10 into a host cell.

12. A recombinant host cell produced by the method of claim 11.

13. A recombinant method for producing a NKX3.1 polypeptide, comprising culturing the recombinant host cell of claim 12 under conditions such that said polypeptide is expressed and recovering said polypeptide.

14. An isolated polypeptide selected from the group consisting of:
(a) a polypeptide comprising amino acids from about 1 to about 234 in SEQ ID NOs:2 or 4;

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- 5 (b) a polypeptide comprising amino acids from about 2 to about 234 in SEQ ID NOs:2 or 4;
- (c) a polypeptide comprising amino acids from about 123 to about 153 in SEQ ID NOs:2 or 4;
- 10 (d) a polypeptide comprising the amino acid sequence encoded by the cDNA clone contained in ATCC Deposit No. 209005 or 209006;
- (e) a polypeptide comprising an epitope bearing portion of any one of the polypeptides of (a), (b), (c), or (d);
- 15 (f) a polypeptide variant created by altering the amino acid sequence of (a), wherein:
- (1) said altering includes an insertion, deletion, or substitution, or any combination thereof; and
- (2) the number of alterations is equal to or less than 5 % of the total number of amino acids present in (a);
- 20 (h) a polypeptide variant created by altering the polynucleotide of (b), wherein:
- (1) said altering includes an insertion, deletion, or substitution, or any combination thereof; and
- (2) the number of alterations is equal to or less than 5 % of the total number of amino acids present in (b);
- 25 (i) a polypeptide variant created by altering the amino acids of (c), wherein:

-109-

- 5
- (1) said altering includes an insertion, deletion, or substitution, or any combination thereof; and
- (2) the number of alterations is equal to or less than 5 % of the total number of amino acids present in (c); and
- (j) a polypeptide variant created by altering the amino acids of (d), wherein:
- 10
- (1) said altering includes an insertion, deletion, or substitution, or any combination thereof; and
- (2) the number of alterations is equal to or less than 5 % of the total number of amino acids present in (d).

15

15. An isolated polypeptide comprising an epitope-bearing portion of the NKX3.1 protein, wherein said portion is selected from the group consisting of: a polypeptide comprising amino acid residues from about 1 to about 13 in SEQ ID NO:2 or SEQ ID NO:4; a polypeptide comprising amino acid residues from about 18 to about 24 in SEQ ID NO:2 or SEQ ID NO:4; a polypeptide comprising amino acid residues from about 35 to about 89 in SEQ ID NO:2 or SEQ ID NO:4; a polypeptide comprising amino acid residues from about 103 to about 113 in SEQ ID NO:2 or SEQ ID NO:4; a polypeptide comprising amino acid residues from about 117 to about 130 in SEQ ID NO:2 or SEQ ID NO:4; and a polypeptide comprising amino acid residues from about 173 to about 185 in SEQ ID NO:2 or SEQ ID NO:4.

20

25

16. An isolated antibody that binds specifically to a NKX3.1 polypeptide of claim 14.

17. An agonist of the polypeptide of claim 14.

18. A method of treating prostate cancer comprising administering to a patient in need thereof an effective amount of the polypeptide of claim 14.

5 19. A method of treating prostate cancer comprising administering to a patient in need thereof an effective amount of the agonist of claim 14.

20. An isolated nucleic acid molecule comprising a polynucleotide encoding a NKX3.1 polypeptide wherein, except for at least one conservative amino acid substitution, said polypeptide has a sequence selected from the group consisting of:

10 (a) a nucleotide sequence encoding a polypeptide comprising amino acids from about 1 to about 234 in SEQ ID NO:2;

(b) a nucleotide sequence encoding a polypeptide comprising amino acids from about 2 to about 234 in SEQ ID NO:2;

15 (c) a nucleotide sequence encoding a polypeptide comprising amino acids from about 1 to about 234 in SEQ ID NO:4;

(d) a nucleotide sequence encoding a polypeptide comprising amino acids from about 2 to about 234 in SEQ ID NO:4;

20 (e) a nucleotide sequence encoding a polypeptide having the amino acid sequence encoded by the clone contained in ATCC Deposit No. 209005 or 209006; and

(f) a nucleotide sequence complementary to any of the nucleotide sequences in (a), (b), (c), (d), or (e).

25 21. An isolated NKX3.1 polypeptide wherein, except for at least one conservative amino acid substitution, said polypeptide has a sequence selected from the group consisting of:

(a) amino acids from about 1 to about 234 in SEQ ID NO:2;

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- (b) amino acids from about 2 to about 234 in SEQ ID NO:2;
(c) amino acids from about 1 to about 234 in SEQ ID NO:4;
(d) amino acids from about 2 to about 234 in SEQ ID NO:4;
(e) the amino acid sequence of the NKX3.1 polypeptide having
5 the amino acid sequence encoded by the clone contained in ATCC Deposit No.
209005 or 209006; and
(f) the amino acid sequence of an epitope-bearing portion of
any one of the polypeptides of (a), (b), (c), (d), or (e).

22. A screening assay for the identification of substances capable of
10 altering the expression from the NKX3.1 promoter, comprising:

- (a) measuring the level of expression of a reporter gene in a
test cell, wherein said test cell is transformed with a recombinant DNA molecule
comprising a reporter gene operably linked to a DNA molecule comprising the
promoter of NKX3.1, and wherein a candidate NKX3.1 *trans*-acting agent is
15 administered to said test cell;
(b) measuring the level of expression of said reporter gene in
a control cell, wherein said control cell is transformed with the recombinant DNA
molecule of step (a); and
(c) comparing the level of expression of said reporter gene in
20 said test cell to the level of said reporter gene in said control cell.

23. A method of determining an animal's predisposition to cancer
comprising screening for a DNA polymorphism at position 154 in SEQ ID NOs:
1 and 3.

24. A prostate-specific gene therapy method comprising
25 (a) introducing a vector containing a gene operably linked to
a NKX3.1 promoter into prostate cells; and
(b) permitting the vector to express the gene in the cells.

25. An isolated nucleic acid molecule comprising nucleotides selected from the group consisting of:

- (a) 50 contiguous nucleotides from the coding region of SEQ ID NO:1; and
- (b) complement of (a).

1 / 15

10 30 50
ATGCTCAGGGTTCCGAGCCGCGGCCCGGGAGGCCAAAGCGAGGGGGCCGCGCCCGC
M L R V P E P R P G E A K A E G A A P P

70 90 110
ACCCCGTCCAAGCCGCTCAGTCCTTCTCATCCAGGACATCCTGCGGGACGGCGCCAG
T P S K P L T S F L I Q D I L R D G A Q

130 150 170
CGGCAAGCGCGCCGACGAGCCAGAGACAGCGGACCCGGAGCCGAGCCAGAGCCA
R Q G G R T S S Q R Q R D P E P E P E P

190 210 230
GAGCCAGAGGGAGGACGAGCCGCGCGGGCGCAGAACGACCAGCTGAGCACCGGGCC
E P E G G R S R A G A Q N D Q L S T G P

250 270 290
CGCGCCGCGCGGAGGAGCGGAGACGCTGGCAGAGACCGAGCCAGAAAGCCTTGGG
R A A P E E A E T L A E T E P E R H L G

310 330 350
TCTTATCTGTTGACTCTGAAAACACTTCAGGCGCCCTTCCAAGCCTTCCCCAAACCCCT
S Y L L D S E N T S G A L P R L P Q T P

370 390 410
AAGCAGCCGAGAACGCTCCCGAGCTGCCTTCTCCCACTCAGGTGATCGAGTTGAG
K Q P Q K R S R A A F S H T Q V I E L E

430 450 470
AGGAAGTTCAGCCATCAGAAGTACCTGTCGGCCCTGAACGGGGCCACCTGGCCAAGAAC
R K F S H Q K Y L S A P E R A H L A K N

490 510 530
CTCAAGCTCAGGAGACCCAAGTGAAGATATGGTTCCAGAACAGACGCTATAAGACTAAG
L K L T E T Q V K I W F Q N R R Y K T K

550 570 590
CGAAAGCAGCTCTCCTCGGAGCTGGGAGACTTGGAGAAGCACTCCTCTTTCGGGCCCTG
R K Q L S S E L G D L E K H S S L P A L

610 630 650
AAAGAGGAGCCTTCTCCCGGCCTCCCTGGTCTCCGTGTATAACAGCTATCCTTACTAC
K E E A F S R A S L V S V Y N S Y P Y Y

670 690
CCATACCTGTACTGCGTGGGCAGCTGGAGCCAGCTTTTGGGTAA
P Y L Y C V G S W S P A F G *

FIG.1

SUBSTITUTE SHEET (RULE 26)

2 / 15

10 30 50
ATGCTCAGGTTCCGGAGCCGGCCCGGGAGCGGAAAGCGGAGGGGCGCGCCGCCG
M L R V P E P R P G E A K A E G A A P P

70 90 110
ACCCCGTCCAAGCCGCTCAGTCTCTCTCATCCAGGACATCCTGCGGAGCGCGCGCAG
T P S K P L T S F L I Q D I L R D G A Q

130 150 170
CGGAAGGGCGCCGACGAGCAGCCAGAGACAGTGGGACCCGAGCCGAGCCAGAGCCA
R Q G G R T S S Q R Q C D P E P E P E P

190 210 230
GAGCCAGAGGGAGGACGACGCCGCGCGGGCGCAGAACGACCAGCTGAGCACCGGGCCC
E P E G G R S R A G A Q N D Q L S T G P

250 270 290
CGCGCCGCGCCGAGGAGCGCGAGACGCTGGCAGAGACCGAGCCAGAAAGGCACTTGGG
R A A P E E A E T L A E T E P E R H L G

310 330 350
TCTTATCTGTTGACTCTGAAAACACTTCAGCGCCCTTCCAAGGCTTCCCCAAACCCCT
S Y L L D S E N T S G A L P R L P Q T P

370 390 410
AAGCAGCCGAGAGCGCTCCCGAGCTGCCCTTCTCCACACTCAGGTGATCGAGTTGGAG
K Q P Q K R S R A A F S H T Q V I E L E

430 450 470
AGGAAGTTCAGCCATCAGAAGTACCTGTCCGCCCTGAACGGGCCACCTGGCCAAGAAC
R K F S H Q K Y L S A P E R A H L A K N

490 510 530
CTCAAGCTCACGGAGACCCAAGTGAAGATATGGTTCCAGAACAGACGCTATAAGACTAAG
L K L T E T Q V K I W F Q N R R Y K T K

550 570 590
CGAAAGCAGCTCTCCTCGGAGCTGGGAGACTTGGAGAAGCACTCTCTTTGCCGGCCCTG
R K Q L S S E L G D L E K H S S L P A L

610 630 650
AAAGAGGAGGCTTCTCCCGGCCCTCCCTGGTCTCCGTGTATAACAGCTATCCTTACTAC
K E E A F S R A S L V S V Y N S Y P Y Y

670 690
CCATACCTGTACTGCGTGGCAGCTGGAGCCAGCTTTTGGGTAA
P Y L Y C V G S W S P A F G *

FIG.2

SUBSTITUTE SHEET (RULE 26)

Figure 3B

Mouse Human	MLRVAEPRPRVETAGGRSPKWAAPPTQSKRLTSFLIQDILRDRAERHSGHGNP-QHSPPRRDSA MLRVPEPRPGEAKEAGAAPPTP-----SKPLTSELIQDILRDRAERHSGHGNP-QHSPPRRDSA	N-TERMINAL REGION 43.1% IDENTITY
Mouse Human	PEPDKAGRGVAPEDPPSIRHSPAEPT-----TEPESDAEHETVLLDCENPGLASAPQVTKQP PEGGRSRAGAQNQLSTGPRAPAEAEETLAETEPER-HIGSVLLDSNTSCALPRLPQTPKQP	43.1% IDENTITY
Mouse Human	QRRSRAAFSHTQVIELERKFSHQKYL SAPERAHAKNLKLTETQVKIWFQNRRYKTKRKQ QRRSRAAFSHTQVIELERKFSHQKYL SAPERAHAKNLKLTETQVKIWFQNRRYKTKRKQ	HOMOBOMAIN 100% IDENTITY
Mouse Human	LSSELGVLEKMSLSLPALKDDSLPSTSELVSVYTSYPYPYLYCLGSMHPSFN LSSELGVLEKMSLSLPALKDDSLPSTSELVSVYTSYPYPYLYCLGSMHPSFN	C-TERMINAL REGION 64.7% IDENTITY

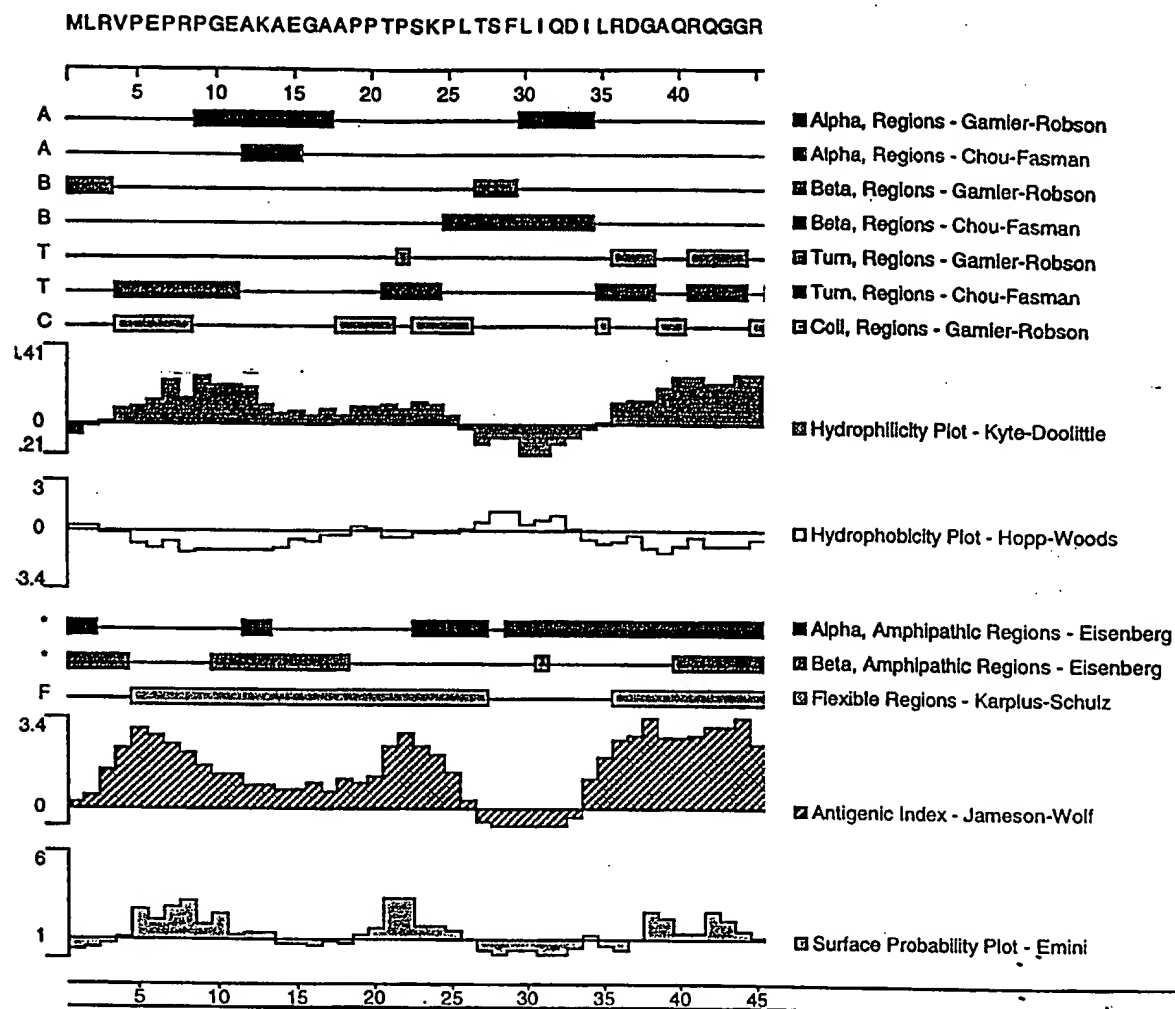


Figure 4

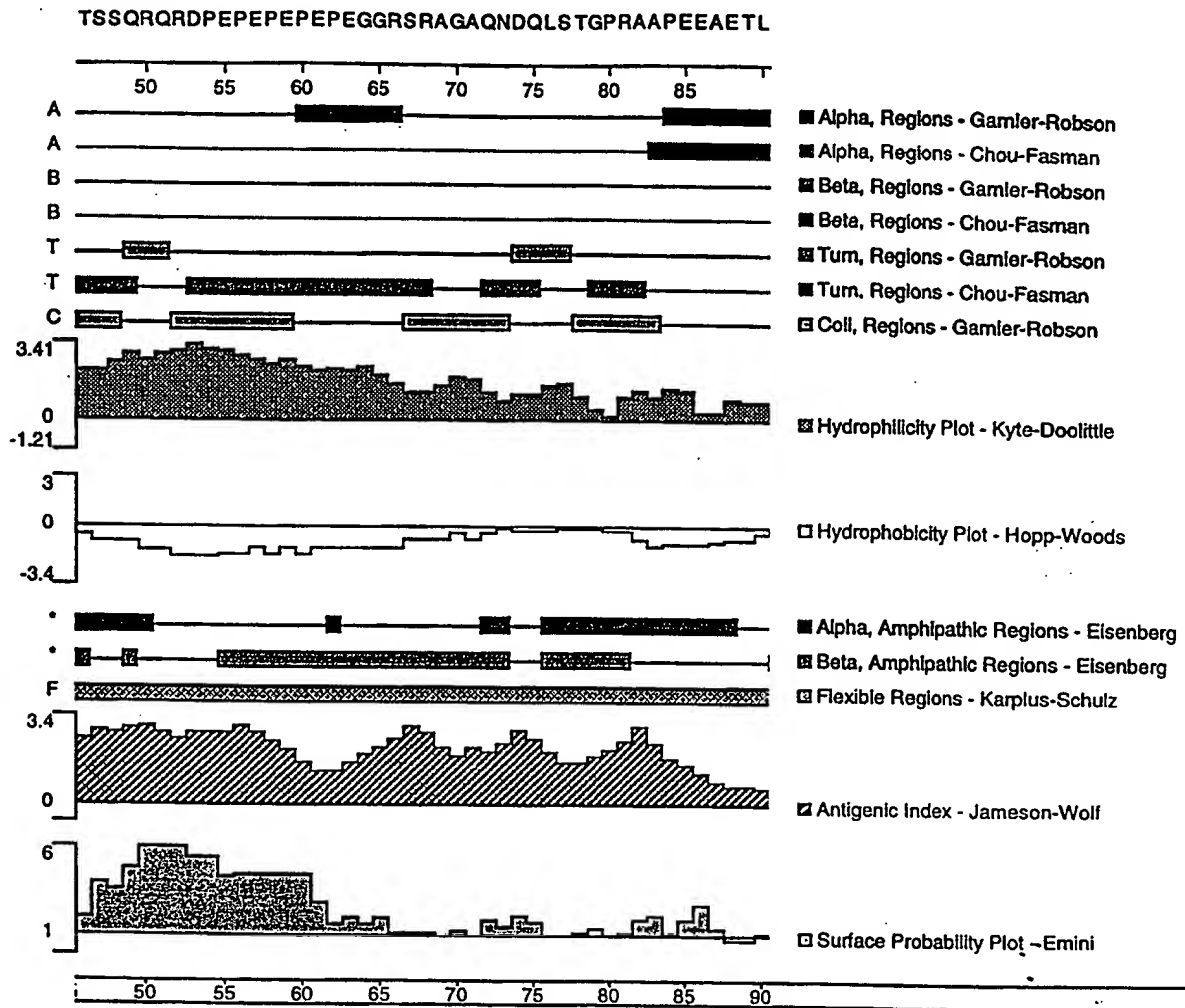
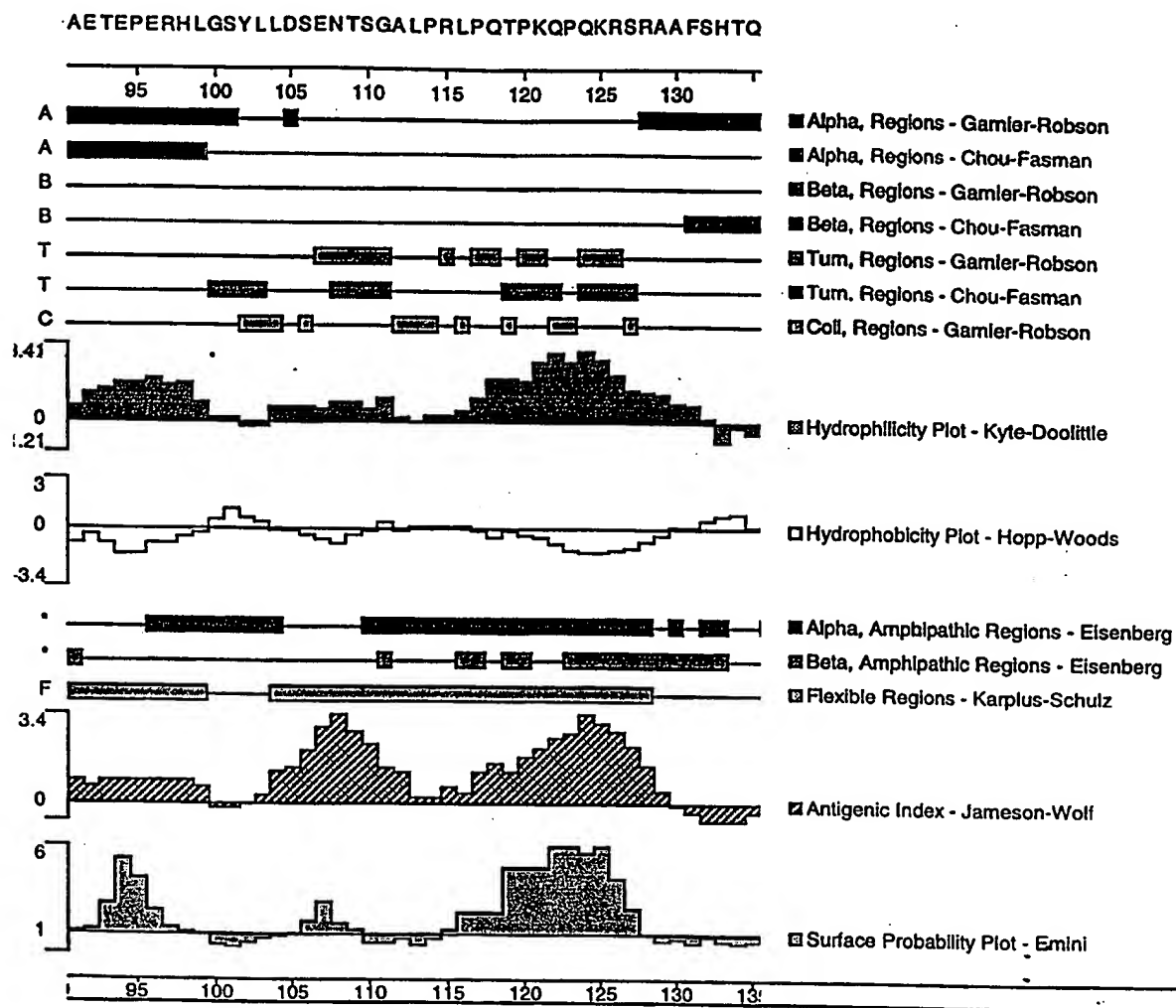


Figure 4 (con't)

*Figure 4 (con't)*

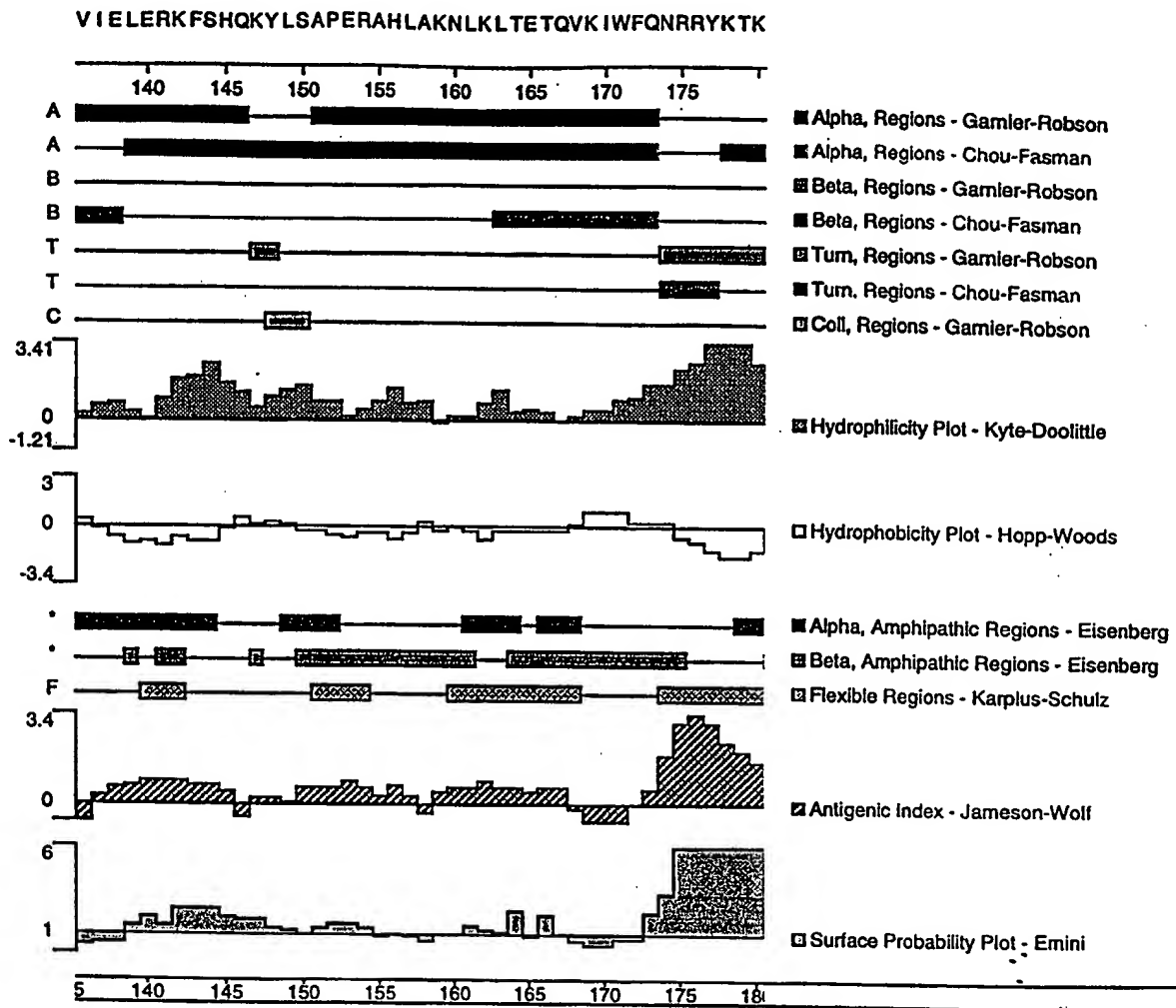


Figure 4 (con't)

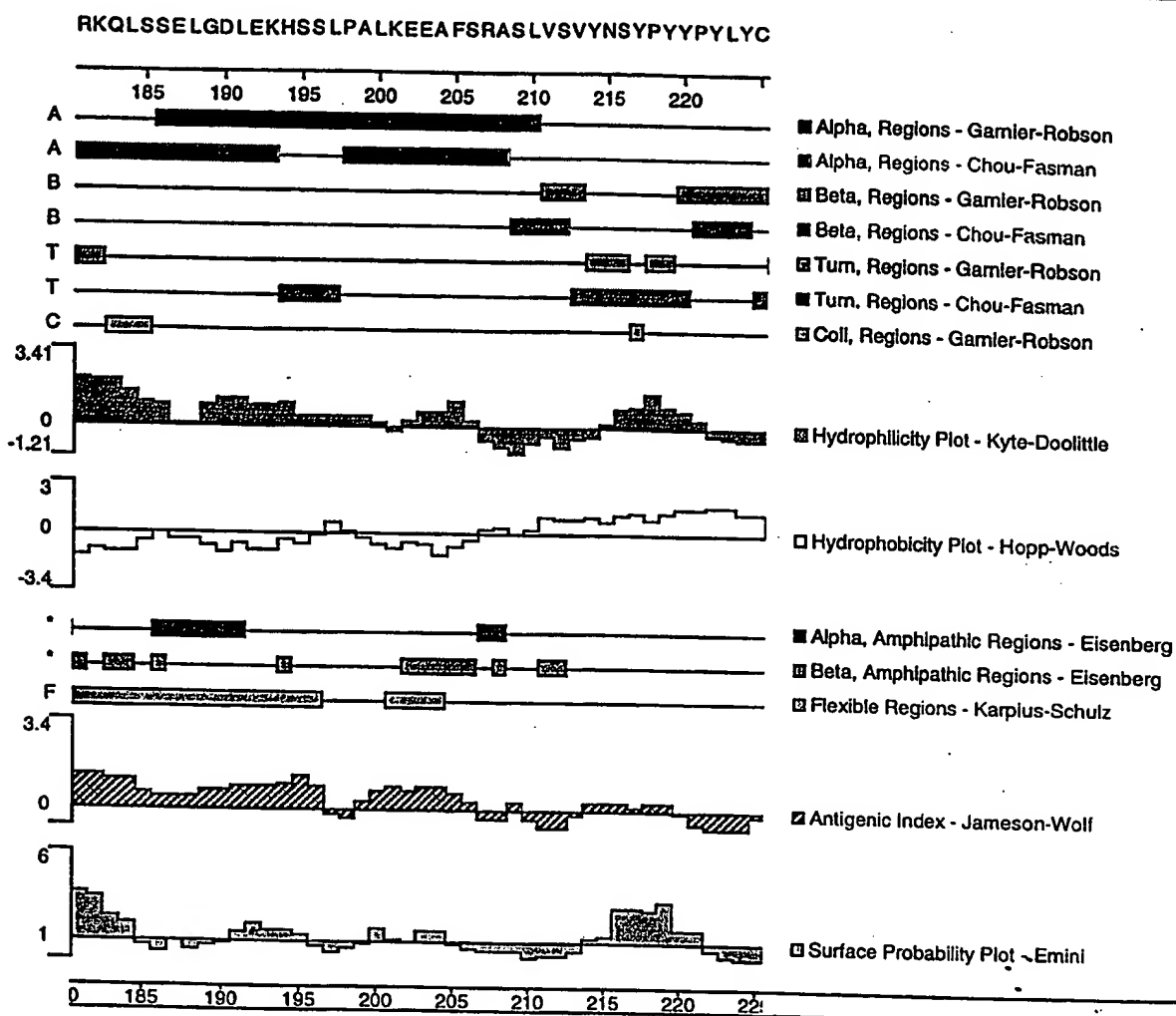


Figure 4 (con't)

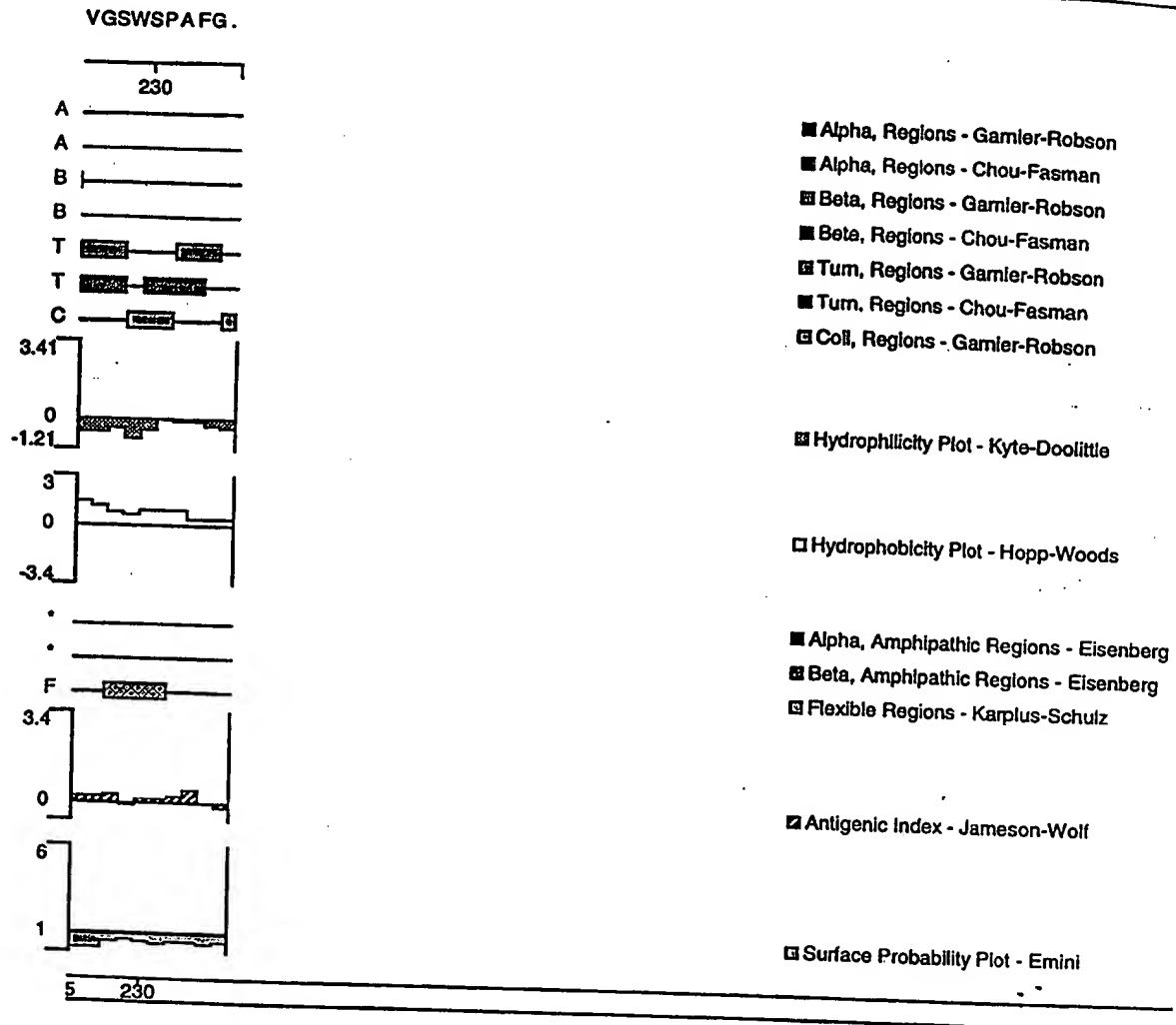
*Figure 4 (con't)*

Figure 5

```

1      AATTAAACCCCT CACTAAAGGG AACAAAAGCT GGAGCTCCAC
41     CGCGGTGGCG GCCGGTAAT ACGACTCACT ATAGGGCGAA
81     GAATTCGGAT CTATCAATCT GCATCCTTGT TTCAGAACCA
121    TTTGATGTAA GTTTCATAAA TCTTGTGCCT TTGCTCCTAC
161    TTTACTTCAGT GTTTATTTCC TAAAAATATT CTCTTGTAACA
201    CTGACAGTAC AATGTGCAAT TTCAGTAAAT TTAACATTAA
241    TTCAATACTT CCATCATCGA CCTGACACTG AGACTCATGC
281    CTGTAGTCCT GGCACTTTGA GAGGCCAAGG CAGGAGGATC
321    ACTTGAATCC AGGAAATCGA GGCTGCAGTG AGTTATGATG
361    GCATCACATGC ACTCCAGCCT GGGCGGCAGA GGGAGACCCCT
401    GTCCGTAAAA AACAGAAGAG AAAAGACAAG GAAAGAAAAT
441    ACTTCCATCA TCTCTGTTC ACCTTCGTCT GTTGTCACGG
481    TACCGTCCAG TCCAGTCACA GTACCGGTTG GACCAATCTG
521    GCTAACCCAT TGTTTAGCCA ATGGGTTACA TGTTAACAGT
561    TGGTAATCTG CAAAAGAGT ATGCTGATGT TCTTTTGAAC
601    TACTTTTTTA AATGCAGTTT TTGCATTTGT CCTGGCCTA
641    AAACGCCCTTC CATCCGTCCTG GAAACTTTTC AAAAGGATGG
681    TATGTTCATGT GTCTGGGGAG GAAGGAAAGT TAACAGGTTA
721    TTGCGGATAA AGGAACCACC AAAGAAAACC ACTTCTGCAA
761    CGGGAAAAGG CTTTGGCAAA GGIGTTTTTC TTCTTTTACG
801    CTGGGGTCTG GCTGCACCTA CTGTTCATGC CTCTTTGAGG
841    TCGTAGATAT TGCAGATCTG AGTTTGCACC ATCTCTCCCA
881    GAGAGAGAGA GCACCCAGAA CTCTCACGGT ACCGCGCGGC
921    TGCAGTGAAT GCGTGTCTAT CCCCTGTAAT TGGCTCTGAC
961    GGTCCTGAAG AGCTAACTGG ACTGTTTGTG TTGATCGTCC
1001   CATCCCCAGG AGCTTCTCTC TGCTGCGGGT GGGTTGGGGC
1041   AGAGGAGCCC CGCTTTGGGG TGCGCTCCTG GCCTGGGAAA
1081   ACGGCTCAGG GCGGAGGGAG GAGAGCTGGA GAAGGAGAGG
1121   AAATTGGGGA AGGAGAGGGA ATTGGGGAAG GAGAGGGAAC
1161   TGGGGAAGGA ATCCCTTAGG GAGGAGCGGA GCGGGGCAGT
1201   GCTCAGGGCT CGCAGATCGG CGGGGTACCC TGGGGCTCAG
1241   GCGGGCAGAT CCGGGGCGCG GCCCGTCCCG CGGCCAATGG
1281   GAGGGCGGCG CGGCCCCGCTC CCTGGGGCTA TAAGCGAGCC
1321   GGGAGGCGGA AAGTGAAAGC GGTCGCGGCC GGGCGGGTGC
1361   ATTCAAGCCA AGGCGGGGCC GCGGGGATGC TCAGGGTTCC
1401   GGAGCCGCGG CCGGGGGAGG CGAAAGCGGA GCGGGCCGCG
1441   CCGCCGACCC CGTCCAAGCC GCTCACGTCC TTCTCATCC
1481   AGGACATCCT GCGGGACGGC GCGCAGCGGC AAGGCGGCCG
1521   CACGAGCAGC CAGAGACAGC GCGACCCGGA GCGGAGCCA
1561   GAGCCAGAGC CAGAGGGAGG ACGCAGCCGC GCGGGGGCGC

```

Figure 5 (con't)

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1601 AGAAGACCA GCTGAGCACC GGGCCCCGCG CCGCGCCGGA
1641 GGAGGCCGAG ACGCTGGCAG AGACCGAGCC AGGTAAGCGG
1681 CGAGGCCGGG GAAGGGGGGC AGCCCCAAGGC GGACCCCCAG
1721 AGCTGGGGGT GCAGGGACGC GGGGCTCCGC GGCGACAGGC
1761 AGAGGGACCT TCCCGCCTCC GCAGCCACGC GCGCGCCCCC
1801 GGAATGAACC CTGAGCCCCA GCGTCAGGGC GGCGCAGGAT
1841 TCTGACACCG CAGGATTGCG CCGGTTCGGT GCCTTCCGTT
1881 CCTTGGGGCT CAGAAGCCGG CCGACTGCA GCGCCACCGC
1921 CTTCACCGT CCCAGGAGCG GATCCCGCCC CCGGCCACCC
1961 GCGATCGGCG CCAGCCCCCC GGTAGTTATG AGAANIAATA
2001 ATAAC TTATT AACAGTGACA AAGCAGGGGT TGACCAGCAA
2041 AGCCTCCGTG TGCTTCCCAA TCCCGTGGGC AGTAAAGCGG
2081 TATATTGGG GTTCCCTCCG GTGTCCAGGA GAGAGAGTCC
2121 ACTPATTTTC TTTCCTGTCA CTCTGATGA GCGACCGAA
2161 CGCCTCGTTT AGCGAAGAGG GAATTAAAGC CCAGAATGAG
2201 CCTGCCCTCG CGTCTCCAGT GGCACAAGCC CTCTCTTGCC
2241 CACCTGGATC CTAACACCGG ATGTCTTTTG GTCTGGCCTT
2281 CCCGGGTATC TTGTTCACG GCATTTTCCC TGCCCTCCCTC
2321 TCCCGCCTCT CCTCAGCACA CAGATCCAGA ATCCCCATAT
2361 AATTCTACTA GACAGTAGGG AGAAAGTTCA ACCACGAAAC
2401 GTCCTTAAC TGGGTTCTT GATGATTCTT AGCAAATGAA
2441 TGCGTAATAA ACATATTTAC TCACTCTTCA CTCCGGAGAG
2481 CTCTTAGTTC ATGTGAAAAA AGTGAAATGT ATCCACGATG
2521 ACAGTGGGCT GTTTGTTCAC TCACTAAAGA GATAAGGGTG
2561 GATTGAATTC TGTTCTCTTC CCTGCTAACA TGTAAC TTTT
2601 GTCTTCCCAT CCTTCTTCC CCACTCTCCT TTCCAGAAAG
2641 GCACTTGGGG TCTTATCTGT TGGACTCTGA AAACACTTCA
2681 GCGGCCCTTC CAAGGCTTCC CCAAACCCCT AAGCAGCCGC
2721 AGAAGCGCTC CCGAGCTGCC TTCTCCACA CTCAGGTGAT
2761 CGAGTTGGAG AGGAAGTTCA GCCATCAGAA GTACCTGTGG
2801 GCCCCTGAAC GGGCCACCT GGOCAAGAAC CTCAGCTCA
2841 CCGAGACCCA AGTGAAGATA TGGTTCCAGA ACAGACGCTA
2881 TAAGACTAAG CGAAAGCAGC TCTCCTCGGA GCTGGGAGAC
2921 TTGGAGAAGC ACTCCTCTTT GCGGCCCCTG AAAGAGGAGG
2961 CCTTCTCCCG GGCTCCCTTG GTCTCCGTGT ATAACAGCTA
3001 TCCTTACTAC CCATACCTGT ACTGCGTGGG CAGCTGGAGC
3041 CCAGCTTTTG GGTAAATGCCA GCTCAGGTGA CAACCATTAT
3081 GATCAAAAAC TGCCTTCCCC AGGGTGTCTC TATGAAAAGC
3121 ACAAGGGGCC AAGGTCAGGG AGCAAGAGGT GTGCACACCA
3161 AAGCTATTGG AGATTTGCGT GGAAATCTCA GATTCTTCAC

```

Figure 5 (con't)

3201 TGGTGAGACA ATGAAACAAC AGAGACAGTG AAAGTTTTAA
3241 TACCTAAGTC ATTCTCCAG TGCATACTGT AGGTCATTTT
3281 TTTTGGTTCT GGCTACCTGT TTGAAGGGGA GAGAGGGAAA
3321 ATCAAGTGGT ATTTTCCAGC ACTTTGTATG ATTTTGGATG
3361 AGTTGTACAC CCAAGGATTC TGTATGCAA CTCATCCTC
3401 CTGTGTCACT GAATATCAAC TCTGAAAGAG CAAACCTAAC
3441 AGGAGAAAGG ACAACCAGGA TGAGGATGTC ACCAACTGAA
3481 TTAAACTC

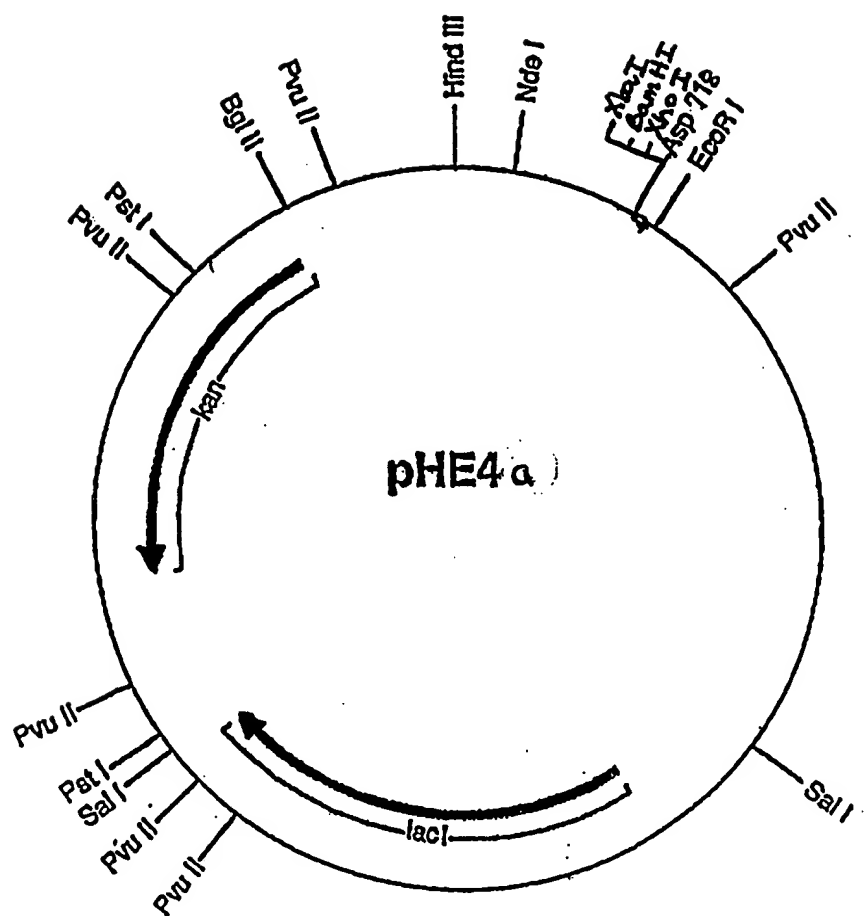


FIG. 6

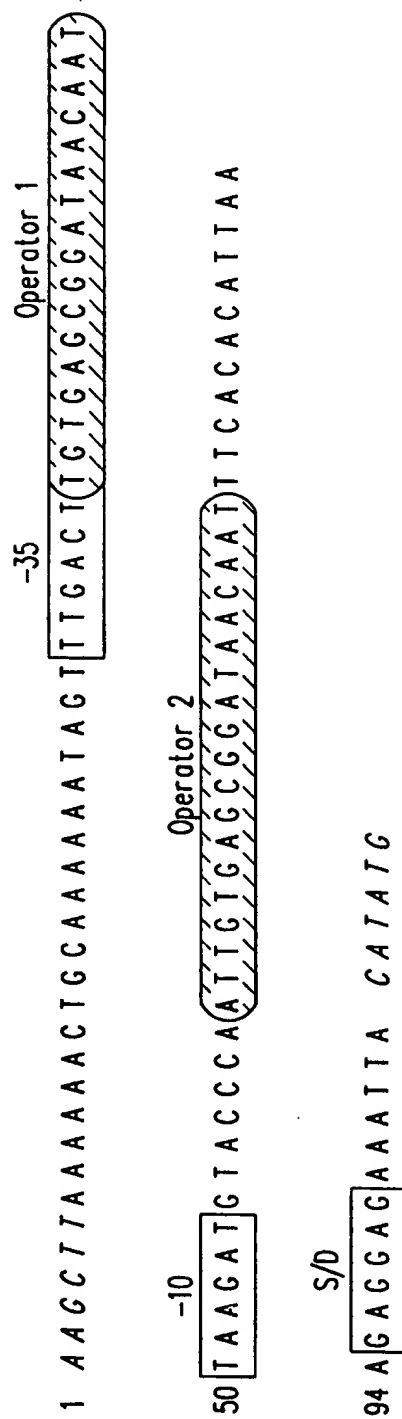


FIG.7

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 98/13252

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C12N15/12 C07K14/47 C12Q1/68 A61K48/00 A61K38/17
C07K16/18

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07K C12N A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	C.J. BIEBERICH ET AL: "Prostate-specific and androgen-dependent expression of a novel homeobox gene" JOURNAL OF BIOLOGICAL CHEMISTRY., vol. 271, no. 50, 13 December 1996, pages 31779-31782, XP002084745 MD US -& EMBL DATABASE ENTRY MMU73460, ACCESSION NUMBER U73460, 16 December 1996, XP002084756 see the whole document --- -/--	1,6-15

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

20 November 1998

Date of mailing of the international search report

04/12/1998

Name and mailing address of the ISA

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Authorized officer

Le Cornec, N

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 98/13252

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	J.L. PRESCOTT ET AL: "human androgen regulated homeobox protein (NKX3.1) mRNA, complete cds" EMBL DATABASE ENTRY HSU80669, ACCESSION NUMBER U80669, 17 December 1996, XP002084746 see abstract ---	1-15,25
X	P.J. SCIAVOLINO ET AL: "Tissue-specific expression of murine NKx3.1 in the male urogenital system" DEVELOPMENTAL DYNAMICS, vol. 209, no. 1, 18 April 1997, pages 127-138, XP002084747 see the whole document ---	1,6-15
A	G.S. BOVA ET AL: "Homozygous deletion and frequent allelic loss of chromosome 8p22 loci in human prostate cancer" CANCER RESEARCH, vol. 53, 1 September 1993, pages 3869-3873, XP002084748 MD US see the whole document ---	23
A	WO 95 32214 A (CANJI, INC.) 30 November 1995 see the whole document ---	
A	WO 97 20942 A (PIAO Y.) 12 June 1997 see abstract; claims: example 9 ---	22
P,X	W.W. HE ET AL: "a novel human prostate-specific androgen-regulated homeobox gene (NKX3.1) that maps to 8p21, a region frequently deleted in prostate cancer" GENOMICS, vol. 43, no. 1, 1 July 1997, pages 69-77, XP002084749 see the whole document ---	1-15,25
P,X	J.L. PRESCOTT ET AL: "Isolation and androgen regulation of the human homeobox cDNA, NKX3.1" THE PROSTATE, vol. 35, no. 1, 1 April 1998, pages 71-80, XP002084750 see the whole document ---	1-15,20, 21,25
A	---	18
	-/--	

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 98/13252

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication where appropriate, of the relevant passages	Relevant to claim No.
P,X	<p>H.J. VOELLER ET AL: "Coding region of NKX3.1, a prostate-specific homeobox gene on 8p21, is not mutated in human prostate cancers"</p> <p>CANCER RESEARCH., vol. 57, 15 October 1997, pages 4455-4459, XP002084751 MD US see the whole document -----</p>	<p>1-13,23, 25</p>

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 98/13252

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
Remark: Although claims 18 and 24 (as far as it concerns an in vivo method) are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☒ Claims Nos.: 17, 19
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:

Claims 17 and 19 are unsearchable because an agonist of the polypeptide NKx3.1 has not been described.
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☒ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International Application No. PCT/ US 98 /13252

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 1-16, 18, 20-21, 23

human NKx3.1. Production by genetic engineering. Antibody.
Pharmaceutical composition.

2. Claims: 22,14

use of the NKX3.1 promoter.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 98/13252

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9532214 A	30-11-1995	AU 686815 B	12-02-1998
		AU 2603495 A	18-12-1995
		CA 2170627 A	30-11-1995
		CZ 9600531 A	14-05-1997
		EP 0714401 A	05-06-1996
		FI 961723 A	19-04-1996
		NO 961539 A	18-04-1996
		SK 24696 A	05-02-1997
WO 9720942 A	12-06-1997	AU 1067697 A	27-06-1997